

**Best  
Available  
Copy**

**AD-A281 193**



1

OFFICE OF NAVAL RESEARCH

CONTRACT N00014-89-J-1828

R&T Code 3132080

Abstract Report #6

POLYMER EFFECT ON HETEROCHIRAL MOLECULAR RECOGNITION IN  
MOLECULAR AND MACROMOLECULAR PAIRS OF LIQUID CRYSTALS OF (R)- AND  
(S)-2-CHLORO-4-METHYLPENTYL 4'-[(8-(VINILOXY)OCTYL)OXY]BIPHENYL-4-  
CARBOXYLATE ENANTIOMERS

**DTIC**  
**S** **ELECTE** **D**  
JUL 11 1994  
**F**

by

V. Percec and H. Oda

Published

in the

**DTIC QUALITY INSPECTED 2**

Macromolecules, in press

Department of Macromolecular Science  
Case Western Reserve University  
Cleveland, OH 44106-7202

June 30, 1994

Reproduction in whole or in part is permitted for any purpose of the United States Government

This document has been approved for public release and sale;  
its distribution is unlimited.

**94-20930**



**94 7 8 03 8**

OMB No. 0704-0184

Operations and Requirements Section  
181 Washington, DC 20531

USA 7540-01-130-5522

**Polymer Effect on Heterochiral Molecular Recognition in Molecular  
and Macromolecular Pairs of Liquid Crystals of (R)- and (S)-2-Chloro-  
4-methylpentyl 4'-(8-Vinyloxyoctyloxy)biphenyl-4-carboxylate  
Enantiomers**

**Virgil Percec\* and Hiroji Oda  
Department of Macromolecular Science  
Case Western Reserve University  
Cleveland, Ohio 44106**

Accession For	
NTIS CRA&I	<input checked="" type="checkbox"/>
DTIC TAB	<input type="checkbox"/>
Unannounced	<input type="checkbox"/>
Justification	
By	
Distribution /	
Availability Codes	
Dist	Avail and/or Special
A-1	

\*To whom all correspondence should be addressed

**Abstract:** (R)-2-Chloro-4-methylpentyl 4'-(8-vinyloxyoctyloxy)biphenyl-4-carboxylate ((R)-8) (R >95%) and (S)-2-chloro-4-methylpentyl 4'-(8-vinyloxyoctyloxy)biphenyl-4-carboxylate ((S)-8) (S >95%) enantiomers and their corresponding homopolymers and copolymers with well defined molecular weight and narrow molecular weight distribution were synthesized and characterized. The phase behavior of the two enantiomeric polymers can be compared only by superimposing the dependence of their transition temperatures as a function of molecular weight. The phase behaviors of (R)-8 and poly[(R)-8] are identical to those of (S)-8 and poly[(S)-8] respectively. Both monomers display monotropic  $S_A$  and  $S_C^*$  phases and a crystalline phase, while the corresponding polymers exhibit enantiotropic  $S_A$ ,  $S_C^*$  and  $S_X$  (unidentified smectic) mesophases. Phase diagrams were investigated in detail in binary mixtures of (R)-8 with (S)-8, poly[(R)-8] with poly[(S)-8] and in binary copolymers of (R)-8 with (S)-8 as a function of the composition of the two enantiomeric structural units. In all these systems the two enantiomeric structural units derived from the two monomers are miscible within all their mesophases and over the entire range of composition. This is in contrast to the crystalline phase of the monomers whose phase diagram displays an eutectic composition. The  $S_A$ -I transition of the binary mixture of (R)-8 with (S)-8 is with 0.4°C higher in the 50/50 mixture than the theoretical value expected for an ideal solution, demonstrating the presence of heterochiral molecular recognition between the two enantiomers in their  $S_A$  phase. Heterochiral recognition was not detected in the  $S_C^*$  phase of any of the enantiomeric pairs investigated. In the polymer mixtures, the chiral recognition observed in the  $S_A$  phase of the monomer mixtures is enhanced with increasing the degree of polymerization (DP) up to about 10 to reach a positive deviation of 1.3°C from the theoretical ideal value. However, polymer mixtures with DP higher than 14 showed smaller positive deviations of their  $S_A$ -I transitions than those of the corresponding monomer mixtures. These results indicate the existence of an optimum molecular weight for the manifestation of heterochiral recognition. In the copolymer system, no clear trend was observed with respect to any phase.

## Introduction

Chiral molecular recognition has recently received an increasing interest in various areas of chemistry.<sup>1</sup> Although a variety of topics are of interest in this field, the most notable series of quantitative studies performed in the last decade refer to the investigation of chiral molecular recognition in monolayers consisting of pairs of enantiomers<sup>2</sup> and diastereomers.<sup>3</sup> The influence of chirality is also of considerable interest in the field of molecular thermotropic liquid crystals. Since the discovery of ferroelectric properties in the chiral smectic C ( $S_C^*$ ) phase, this field became extremely active.<sup>4</sup> In most cases, attention has been paid mainly to the electro-optic properties of chiral liquid crystals. There is a limited number of publications in which the influence of chirality on the phase behavior of low molar mass liquid crystals was investigated as a function of their optical purity.<sup>1a,5-9</sup> In few cases, it has been observed that chiral recognition occurs specifically in layered liquid crystalline phases and as a consequence increases the phase transition temperatures of the racemic mixture by comparison to those of the pure enantiomers.<sup>1a,5,7,8</sup> However, the relationship between molecular structure and the mechanism responsible for the manifestation of heterochiral recognition is far from being elucidated.

We reported a series of experiments on the molecular engineering of side chain liquid crystalline polymers by using living cationic polymerization technique as a tool to generate well defined polymers.<sup>10,11</sup> As part of our continuing efforts to elucidate the molecular-supramolecular structures-properties dependence by this technique, we have initiated a series of systematic investigations on chiral molecular recognition using well defined polymers obtained by the living cationic polymerization. In the first publication on this topic, we reported the observation of heterochiral recognition in molecular and macromolecular pairs of diastereomeric liquid crystals based on (2R, 3S)- and (2S, 3S)-2-fluoro-3-methylpentyl 4'-(11-vinyloxyundecanyloxy)biphenyl-4-carboxylate (**9**).<sup>12</sup> Heterochiral recognition was observed in the  $S_A$  but not in the  $S_C^*$ ,  $S_X$  and crystalline phases of the monomer. Heterochiral recognition was not observed in polymer mixtures and in copolymers.<sup>12</sup>

The first goal of this paper is to describe the synthesis and the living cationic polymerization of (R)-2-chloro-4-methylpentyl 4'-(8-vinyloxyoctyloxy)biphenyl-4-carboxylate ((R)-8) and (S)-2-chloro-4-methylpentyl 4'-(8-vinyloxyoctyloxy)biphenyl-4-carboxylate ((S)-8) enantiomers. The second goal of this paper is to compare the mesomorphic behavior of these two enantiomeric structural units and to investigate the heterochiral recognition in binary mixtures of monomers and of polymers, as well as in copolymers. To our knowledge, this paper will report the first example of heterochiral molecular recognition present in both pairs of monomers and of side chain liquid crystalline polymers.

## **Experimental Section**

### **Materials**

L-Leucine [(S)-(+)-2-amino-4-methylpentanoic acid, 99%, Aldrich], D-leucine [(R)-(-)-2-amino-4-methylpentanoic acid, 99%, Aldrich] and borane-tetrahydrofuran complex (BH<sub>3</sub>·THF, 1.0M solution in THF, Aldrich) were used as received.

Pyridine was heated overnight at 100°C over KOH, distilled from KOH, and then stored over KOH. Dimethyl sulfoxide (DMSO) was heated overnight at 100°C over CaH<sub>2</sub>, distilled from CaH<sub>2</sub> under vacuum, and stored over molecular sieves (4Å). THF was refluxed over LiAlH<sub>4</sub> for several days and distilled from LiAlH<sub>4</sub>. Acetone was stored over anhydrous K<sub>2</sub>CO<sub>3</sub> for several days, filtered and distilled.

CH<sub>2</sub>Cl<sub>2</sub> used as polymerization solvent was first washed with concentrated H<sub>2</sub>SO<sub>4</sub> several times until the acid layer remained colorless, then washed with water, dried over MgSO<sub>4</sub>, refluxed over CaH<sub>2</sub>, and freshly distilled under argon before each use. Dimethyl sulfide [(CH<sub>3</sub>)<sub>2</sub>S] used in polymerizations (Aldrich, anhydrous, 99+%, packed under nitrogen in Sure/Seal™ bottle) was used as received. Trifluoromethanesulfonic acid (CF<sub>3</sub>SO<sub>3</sub>H) used as polymerization initiator (Aldrich, 98%) was distilled under vacuum.

All other materials were commercially available and were used as received.

### **Techniques**

<sup>1</sup>H-NMR spectra were recorded on Varian XL-200 (200 MHz) spectrometer.

Relative molecular weights of polymers were determined by gel permeation chromatography (GPC). GPC analyses were carried out with a Perkin-Elmer Series 10LC instrument equipped with an LC-100 column oven and a Nelson Analytical 900 Series data station. Measurements were made by using a UV detector, THF as solvent (1ml/min, 40°C), a set of PL gel columns of  $5 \times 10^2$  and  $10^4 \text{ \AA}$ , and a calibration plot constructed with polystyrene standards.

A Perkin-Elmer PC Series DSC-7 differential scanning calorimeter was used to determine the thermal transition temperatures, which were reported as the maxima and minima of their endothermic or exothermic peaks respectively. Heating and cooling rates were 20°C/min for the analysis of the homopolymers and 10°C/min for the analysis of the monomer and polymer mixtures and of copolymers.

A Carl-Zeiss optical polarizing microscope equipped with a Mettler FP-82 hot stage and a Mettler FP-80 central processor was used to observe the thermal transitions and to analyze the anisotropic textures.

### Synthesis of monomers

Monomers (R)-8 and (S)-8 were synthesized from D-(R)-leucine and L-(S)-leucine respectively according to the reaction pathway outlined in Scheme 1. The synthesis of compounds 5 and 7 was described previously.<sup>11</sup> This synthetic procedure will be described in detail only for monomer (S)-8.

#### (S)-2-Chloro-4-methylpentanoic Acid [(S)-2]<sup>13</sup>

A solution of  $\text{NaNO}_2$  (19.7g, 0.286mol) in water (50ml) was added dropwise to a stirred, cooled (0°C) mixture of L-leucine (25.0g, 0.191mol), concentrated HCl (125ml) and water (125ml) over 1.5 h. Stirring was continued overnight while the mixture was allowed to warm to room temperature. The product was extracted into diethyl ether four times and the combined ethereal extracts were dried over anhydrous  $\text{MgSO}_4$ . The solvent was evaporated and the remaining crude product was distilled under vacuum to yield a colorless liquid (18.4g,



64.1%). Bp 82-86°C (2mmHg).  $^1\text{H-NMR}$  ( $\text{CDCl}_3$ , TMS):  $\delta$  1.00 (m, 6H,  $(\text{CH}_3)_2\text{CH-}$ ), 1.88 (m, 3H,  $(\text{CH}_3)_2\text{CHCH}_2\text{-}$ ), 4.36 (m, 1H,  $-\text{CHCl-}$ ), 10.23 (bs, 1H,  $-\text{COOH}$ ).

**(S)-2-Chloro-4-methylpentanol [(S)-3]**

A solution of  $\text{BH}_3\text{-THF}$  complex in THF (1.0M, 62.5ml, 62.5mmol) was added dropwise to a stirred, cooled (0°C) solution of (S)-2 (7.84g, 52.1mmol) in dry THF (150ml) over 1 hour under nitrogen atmosphere. The reaction mixture was allowed to warm to room temperature and stirring was continued for 2 h. Then the reaction mixture was cooled to 0°C again and 7ml of water followed by 150ml of saturated  $\text{K}_2\text{CO}_3$  solution were added. The product was extracted into diethyl ether twice and the combined organic layers were dried over anhydrous  $\text{MgSO}_4$ . The solvent was evaporated and the remaining crude product (6.43g, 90.3%) was used for the synthesis of (S)-4 without further purification.  $^1\text{H-NMR}$  ( $\text{CDCl}_3$ , TMS):  $\delta$  0.92 (d,  $J=6.5\text{Hz}$ , 3H,  $(\text{CH}_3)_2\text{CH-}$ ), 0.96 (d,  $J=6.5\text{Hz}$ , 3H,  $(\text{CH}_3)_2\text{CH-}$ ), 1.39-1.58, 1.58-1.79 (m, 2H,  $(\text{CH}_3)_2\text{CHCH}_2\text{-}$ ), 1.79-1.97 (m, 1H,  $(\text{CH}_3)_2\text{CHCH}_2\text{-}$ ), 2.19 (bs, 1H,  $-\text{CH}_2\text{OH}$ ), 3.57-3.89 (m, 2H,  $-\text{CH}_2\text{OH}$ ), 4.11 (m, 1H,  $-\text{CHCl-}$ ).

**(S)-2-Chloro-4-methylpentyl Tosylate [(S)-4]**

A solution of (S)-3 (6.43g, 47.1mmol, crude) in dry pyridine (20ml) was added dropwise to a stirred, cooled (0°C) solution of p-toluenesulfonyl chloride (13.5g, 70.7mmol) in dry pyridine (80ml). The mixture was allowed to warm to room temperature and stirred overnight. The mixture was poured into water and the product was extracted into diethyl ether twice. The combined ethereal extracts were washed with 10%  $\text{HCl}$  twice and dried over anhydrous  $\text{MgSO}_4$ . The solvent was evaporated and the remaining crude product was purified by column chromatography (silica gel; hexane-ethyl acetate 15:1) to give a colorless liquid (10.2g, 74.5%). Purity: >99% (TLC).  $^1\text{H-NMR}$  ( $\text{CDCl}_3$ , TMS):  $\delta$  0.86 (d,  $J=6.5\text{Hz}$ , 3H,  $(\text{CH}_3)_2\text{CH-}$ ), 0.92 (d,  $J=6.5\text{Hz}$ , 3H,  $(\text{CH}_3)_2\text{CH-}$ ), 1.50-1.61 (m, 2H,  $(\text{CH}_3)_2\text{CHCH}_2\text{-}$ ), 1.78-1.94 (m, 1H,  $(\text{CH}_3)_2\text{CH-}$ ), 2.46 (s, 3H,  $-\text{PhCH}_3$ ), 3.97-4.18 (m, 3H,  $-\text{CHClCH}_2\text{O-SO}_2\text{-}$ ), 7.37 (d,  $J=8.2\text{Hz}$ , 2ArH, ortho to  $-\text{CH}_3$ ), 7.82 (d,  $J=8.2\text{Hz}$ , 2ArH, ortho to  $-\text{SO}_2\text{-}$ ).

**(S)-2-Chloro-4-methylpentyl 4'-Hydroxybiphenyl-4-carboxylate [(S)-6]**

A mixture of (S)-4 (8.72g, 30.0mmol), 5 (7.57g, 30.0mmol), tetrabutylammonium hydrogen sulfate (TBAH, 1.50g) and dry DMSO (100ml) was stirred at 80°C under nitrogen atmosphere for 20 h. The resulting clear yellow solution was poured into water. The product was extracted into diethyl ether twice and the combined ethereal extracts were dried over anhydrous MgSO<sub>4</sub>. The solvent was evaporated and the remaining crude product was purified by column chromatography twice (silica gel; hexane-ethyl acetate 3:1) to give a white solid (7.00g, 73.6%). Purity: >99% (TLC). Mp 81.2°C (DSC, 20°C/min). <sup>1</sup>H-NMR (CDCl<sub>3</sub>, TMS): δ 0.96 (d, J=6.6Hz, 3H, (CH<sub>3</sub>)<sub>2</sub>CH-), 0.99 (d, J=6.6Hz, 3H, (CH<sub>3</sub>)<sub>2</sub>CH-), 1.52-1.87 (m, 2H, (CH<sub>3</sub>)<sub>2</sub>CHCH<sub>2</sub>-), 1.87-2.11, (m, 1H, (CH<sub>3</sub>)<sub>2</sub>CH-), 4.19-4.59 (m, 3H, -CHClCH<sub>2</sub>OCO-), 5.28 (s, 1H, -PhOH), 6.95 (d, J=8.7Hz, 2ArH, ortho to -OH), 7.54 (d, J=8.7Hz, 2ArH, meta to -OH), 7.63 (d, J=8.5Hz, 2ArH, meta to -COO-), 8.12 (d, J=8.5Hz, 2ArH, ortho to -COO-).

**(S)-2-Chloro-4-methylpentyl 4'-(8-Vinyloxyoctyloxy)biphenyl-4-carboxylate [(S)-8]**

A mixture of (S)-6 (3.50g, 10.5mmol), anhydrous K<sub>2</sub>CO<sub>3</sub> (3.80g, 27.5mmol) and dry acetone (90ml) was stirred at 60°C under a nitrogen atmosphere for 2 h. To the resulting yellow solution was added a solution of 7 (2.59g, 11.0mmol) in dry DMSO (5.0ml) and stirring was continued at 60°C for 20 h. The mixture was poured into water and the product was extracted into diethyl ether twice. The combined ethereal extracts were dried over anhydrous MgSO<sub>4</sub>. The solvent was evaporated and the remaining crude product was purified by column chromatography twice (silica gel; hexane-ethyl acetate 20:1) to yield a white solid (3.20g, 62.6%). Purity: >99% (TLC). The thermal transition temperatures are given in Table IV. <sup>1</sup>H-NMR (CDCl<sub>3</sub>, TMS): δ 0.96 (d, J=6.6Hz, 3H, (CH<sub>3</sub>)<sub>2</sub>CH-), 0.99 (d, J=6.6Hz, 3H, (CH<sub>3</sub>)<sub>2</sub>CH-), 1.18-2.06 (m, 15H, (CH<sub>3</sub>)<sub>2</sub>CHCH<sub>2</sub>-, CH<sub>2</sub>=CHOCH<sub>2</sub>(CH<sub>2</sub>)<sub>6</sub>CH<sub>2</sub>O-), 3.68 (t, J=6.5Hz, 2H, CH<sub>2</sub>=CHOCH<sub>2</sub>-), 3.98 (dd, J=6.8, 1.8Hz, 1H, CH<sub>2</sub>=CHO- trans), 4.01 (t, J=6.5Hz, 2H, -CH<sub>2</sub>O-), 4.18 (dd, J=14.4, 1.8Hz, 1H, CH<sub>2</sub>=CHO- cis), 4.22-4.57 (m,

3H,  $-\underline{\text{CH}}\text{ClCH}_2\text{OCO}-$ ), 6.48 (dd,  $J=14.4, 6.8\text{ Hz}$ , 1H,  $\text{CH}_2=\underline{\text{CH}}\text{O}-$ ), 6.99 (d,  $J=8.7\text{ Hz}$ , 2ArH, ortho to  $-(\text{CH}_2)_8\text{O}-$ ), 7.57 (d,  $J=8.7\text{ Hz}$ , 2ArH, meta to  $-(\text{CH}_2)_8\text{O}-$ ), 7.65 (d,  $J=8.3\text{ Hz}$ , 2ArH, meta to  $-\text{COO}-$ ), 8.11 (d,  $J=8.3\text{ Hz}$ , 2ArH, ortho to  $-\text{COO}-$ ).

### Cationic Polymerization

Polymerizations and copolymerizations were carried out in a three-necked round bottom flask equipped with a Teflon stopcock and rubber septa under argon atmosphere at  $0^\circ\text{C}$  for 1 hour. All glassware was dried overnight at  $140^\circ\text{C}$ . The monomer was further dried under vacuum overnight in the polymerization flask. After the flask was filled with argon, freshly distilled dry methylene chloride was added via a syringe and the solution was cooled to  $0^\circ\text{C}$ . Dimethyl sulfide and trifluoromethanesulfonic acid were then added carefully via a syringe. The monomer concentration was about 0.224 M and the dimethyl sulfide concentration was 10 times larger than that of trifluoromethanesulfonic acid. The polymer molecular weight was controlled by the monomer/initiator ( $[\text{M}]_0/[\text{I}]_0$ ) ratio. After quenching the polymerization with a mixture of  $\text{NH}_4\text{OH}$  and methanol (1:2), the reaction mixture was poured into methanol to give a white precipitate. The obtained polymer was purified by the reprecipitation by pouring its chloroform solution into methanol, and dried under vacuum.

### Results and Discussion

#### Determination of the optical purities of monomers (R)-8 and (S)-8

The synthesis of the two enantiomeric monomers (R)- and (S)-8 is outlined in Scheme 1. Monomers (R)-8 and (S)-8 were synthesized from D-(R)-leucine and L-(S)-leucine respectively in the same manner. The amino groups of D-(R)- and L-(S)-leucine were substituted with a chlorine atom via a diazonium salt by using  $\text{NaNO}_2$  in 6N HCl. Since this reaction proceeds with the retention of configuration at the chiral center, the original configurations of D-(R)- and L-(S)-leucine are maintained throughout the entire steps of the synthesis. The reduction of the carboxyl groups of compounds (R)- and (S)-2 was carried out with a  $\text{BH}_3\text{-THF}$  complex without affecting the chlorine atom of the chiral center.

Reduction experiments with  $\text{LiAlH}_4$  showed that it reduced the chlorine atom, resulting in the partial loss of the chiral center.

The optical purities of monomers (R)- and (S)-8, and of the corresponding polymers poly[(R)-8] and poly[(S)-8] were investigated by the NMR chiral shift reagent technique described by Goodby et al.<sup>7b,14</sup> The chiral shift reagent used in this study was tris[3-(heptafluoro-propylhydroxymethylene)-(+)-camphorato] europium(III) derivative  $[\text{Eu}(\text{hfc})_3]$ . Figure 1 presents the  $^1\text{H}$ -NMR spectra of monomer (R)-8 and polymer poly[(R)-8] (DP=5.7) both without  $\text{Eu}(\text{hfc})_3$ . Figure 2a presents the  $^1\text{H}$ -NMR spectrum of the 50/50 mixture between (R)- and (S)-8 with  $\text{Eu}(\text{hfc})_3$  [(R)-8 5mg, (S)-8 5mg and  $\text{Eu}(\text{hfc})_3$  10mg]. The  $^1\text{H}$ -NMR spectra of pure (R)- and (S)-8 with  $\text{Eu}(\text{hfc})_3$  [monomer 10mg and  $\text{Eu}(\text{hfc})_3$  10mg] are presented in Figures 2b and 2c respectively. As shown in Figure 2, the doublet associated with the aromatic proton j which is adjacent to the chiral center is split in the 50/50 mixture of the monomers, while the corresponding peaks remained unchanged in pure (R)-8 and (S)-8 except that they were shifted down field. When the amount of the shift reagent was increased, a broadening of the peaks was observed before the complete separation of the two doublets was achieved and therefore, it was difficult to calculate the optical purities of both monomers precisely. However, judging from Figures 2b and 2c, it seems that no racemization has occurred throughout the synthesis of the monomers and their optical purities are expected to be higher than 95%. The same technique was also applied to polymers. The polymers showed broadened peaks even in the absence of the shift reagent (Figure 1b) and this made the separation of the peaks impossible. However, it is believed that the chiral center of the monomers is insensitive to the cationic polymerization condition and that the original optical purities of the monomers remain unchanged during this polymerization process. This was the case in previous examples of cationic polymerization of similar chiral monomers with the same initiating system.

## Homopolymerization of (R)-8 and (S)-8

The homopolymerizations of (R)-8 and (S)-8 are presented in Scheme 2. All polymerizations were carried out at 0°C in CH<sub>2</sub>Cl<sub>2</sub> by a living cationic polymerization technique using CF<sub>3</sub>SO<sub>3</sub>H/(CH<sub>3</sub>)<sub>2</sub>S as an initiation system. Previous work in our laboratory<sup>15</sup> and others<sup>16</sup> has shown that the CF<sub>3</sub>SO<sub>3</sub>H initiated polymerization of vinyl ethers in the presence of a Lewis base such as (CH<sub>3</sub>)<sub>2</sub>S gives well defined polymers with controlled molecular weights and narrow polydispersities. The polymerization mechanism is discussed in detail in previous publications.<sup>10,11,15</sup>

The characterization of poly[(R)-8] and poly[(S)-8] by gel permeation chromatography (GPC) and differential scanning calorimetry (DSC) is summarized in Tables I and II respectively. The low polymer yields are the result of the loss of polymer during purification. Relative number-average molecular weights of polymers determined by GPC exhibit a linear dependence on the initial molar ratio of monomer to initiator ( $[M]_0/[I]_0$ ) as shown in Figure 3. All polydispersities are less than 1.23. The  $[M]_0/[I]_0$  ratio provides a good control of the polymer molecular weight. All these features demonstrate the typical characteristics of the living polymerization mechanism. The absolute number-average molecular weights were difficult to determine by <sup>1</sup>H-NMR spectroscopy from the chain ends of the polymer owing to signal overlap.

The mesomorphic behaviors of poly[(R)-8] and poly[(S)-8] were investigated by DSC and thermal optical polarized microscopy. Figures 4 and 5 present the DSC thermograms of poly[(R)-8] and poly[(S)-8] with various degrees of polymerization (DP) respectively. The phase behaviors of the two enantiomeric homopolymers can be compared by superimposing the plots of the dependencies of their thermal transition temperatures as a function of DP (Figure 6). As observed from this figure, the phase behavior of poly[(R)-8] is identical to that of poly[(S)-8]. The DSC curves of first heating scans are very similar to those of second heating scans except that medium molecular weight polymers (DP=6.9-8.7) exhibit a small crystalline melting peak on the first heating scan. On the second and subsequent

heating scans, all polymers show a  $S_C^*$  phase followed by a  $S_A$  phase which melts into an isotropic phase. In polymers with  $DP > 7$ , another higher order smectic phase ( $S_X$ ) is observed below the  $S_C^*$  phase. The nature of this  $S_X$  phase was not identified. The  $S_X$  and  $S_C^*$  phases are enantiotropic phases since these phases are also observed in first and subsequent cooling scans. The  $S_A$  phase, however, is not observed during the cooling scans. This is most probably due to the thermal instability of the  $S_A$  phase whose temperature range is so narrow that the transitions are overlapped with  $I-S_C^*$  transitions under the relatively high rate of the DSC ( $20^\circ\text{C}/\text{min}$ ). In fact, DSC scans at lower cooling rate (e.g.  $10^\circ\text{C}/\text{min}$ ) give rise to the very narrow  $S_A$  phase in between the  $I$  and  $S_C^*$  phases. Representative optical micrographs of the texture exhibited by the  $S_A$  phase of poly[(S)-8] ( $DP=11.8$ ) are presented in Figure 7.

Since no definitive evidence for the existence of the  $S_C^*$  phase (e.g. a focal conic texture with equidistant lines) was obtained from the optical microscope analysis, copolymerization experiments between monomer (S)-8 and (2S, 3S)-2-fluoro-3-methylpentyl 4'-(11-vinyloxyundecanyloxy)biphenyl-4-carboxylate ((2S,3S)-9), whose homopolymer displays  $S_A$  and  $S_C^*$  phases,<sup>12</sup> were performed to confirm further the phase assignment. Based on our previous experience, copolymers derived from two monomers which lead to two homopolymers displaying the same mesophase will show a continuous dependence of their phase transitions on copolymer composition. On the contrary, if the structural units of these two homopolymers are not isomorphic within a particular mesophase, a triple point will occur on the phase diagram.<sup>11b-d, 17</sup> Scheme 3 illustrates the copolymerization of (S)-8 with (2S, 3S)-9. Attempts were made to synthesize poly[[(S)-8]]-co-[(2S,3S)-9]]X/Y (where X/Y refers to the mole ratio of the two structural units) with  $DP=20$ . The copolymerization results are summarized in Table III. The copolymer yields are lower than quantitative due to the polymer losses during the purification process. However, all conversions were quantitative and therefore, the copolymer composition is identical to that of the monomer feed.<sup>10</sup> The DSC traces and the phase diagrams of the copolymers are presented in Figures 8 and 9 respectively. It is clear from these figures that the  $S_A$  and  $S_C^*$  mesophases display a continuous dependence

over the entire composition range, supporting the above phase assignment of poly[(S)-8]. It also appears that the two  $S_X$  phases of poly[(S)-8] and poly[(2S,3S)-9] are isomorphic.

### Miscibility Studies

Monomer (R)-8 and monomer (S)-8 were mixed in various compositions and the phase behavior of their mixtures was investigated by DSC. Mixtures were prepared by dissolving the two monomers in  $\text{CH}_2\text{Cl}_2$  followed by evaporation of the solvent under vacuum. Four sets of binary mixtures between poly[(R)-8] and poly[(S)-8] with different molecular weights were also prepared and their phase behaviors were investigated in the same manner. The molecular weights and polydispersities of the polymers employed in this miscibility study are as follows: Polymer mixture I: poly[(R)-8] with  $\text{DP}=5.7$ ,  $M_w/M_n=1.14$  and poly[(S)-8] with  $\text{DP}=6.1$ ,  $M_w/M_n=1.23$ ; Polymer mixture II: poly[(R)-8] with  $\text{DP}=6.7$ ,  $M_w/M_n=1.23$  and poly[(S)-8] with  $\text{DP}=6.9$ ,  $M_w/M_n=1.21$ ; Polymer mixture III: poly[(R)-8] with  $\text{DP}=9.6$ ,  $M_w/M_n=1.16$  and poly[(S)-8] with  $\text{DP}=10.1$ ,  $M_w/M_n=1.16$ ; Polymer mixture IV: poly[(R)-8] with  $\text{DP}=15.1$ ,  $M_w/M_n=1.20$  and poly[(S)-8] with  $\text{DP}=14.3$ ,  $M_w/M_n=1.19$ . The thermal transition temperatures and the corresponding enthalpy changes are summarized in Table IV (for the monomer mixtures) and Tables V-VIII (for the polymer mixtures). The phase diagrams of the monomer mixtures are presented in Figure 10 and the phase diagrams of the polymer mixtures are presented in Figures 11-14. The DSC thermograms of the monomer mixtures and Polymer mixture III are presented in Figures 15 and 16 respectively.

The phase behavior of monomer (R)-8 is exactly identical to that of monomer (S)-8 and both monomers display monotropic  $S_A$  and  $S_C^*$  phases and a crystalline phase (Figure 10). When the two monomers were mixed, DSC heating scans showed two eutectic points, while on coolings a large crystalline phase suppression with one eutectic point was observed. On the second and subsequent heating scans, the two pure monomers and a 50/50 monomer mixture showed a crystallization peak. In the  $S_A$  and  $S_C^*$  phases obtained during the cooling scans, the two enantiomeric structural units of the monomers are miscible and isomorphic over the entire range of compositions. The  $S_A$ -I transition temperatures showed an upward

curvature, i.e., a positive deviation ( $0.4^{\circ}\text{C}$ ) from the linear dependence predicted by the Schröder-van Laar equation for an ideal solution<sup>18,19</sup>, demonstrating the presence of the chiral molecular recognition between the two enantiomeric structural units. On the contrary, the  $S_C^*-S_A$  transition temperatures showed negative deviations ( $-0.5^{\circ}\text{C}$ ) from the theoretical values. These two contrasting behaviors shown by the  $S_A$ -I and  $S_C^*-S_A$  transitions are identical to those reported in a previous publication on the phase behavior of two diastereomeric liquid crystalline monomer pairs.<sup>12</sup>

In the four sets of polymer mixtures with different molecular weights, the two enantiomeric structural units derived from the two monomers are also miscible and isomorphic in all mesophases across the full composition range. The  $S_A$ -I transitions always show an upward curvature on the second heating scans and the first cooling scans.

In order to compare the  $S_A$ -I transition temperatures quantitatively, all the  $S_A$ -I transition temperatures obtained from the four sets of polymer mixtures are summarized in Figure 17 (open symbols) together with the corresponding theoretical values calculated from the Schröder-van Laar equation (closed symbols). Since the enthalpy changes associated with the  $S_A$ -I transitions are almost identical for the two enantiomeric polymers possessing similar molecular weights, the Schröder-van Laar equation predicts for an ideal solution like mixture a linear dependence of the  $S_A$ -I transition temperatures on composition in all cases. Consequently, the upward curvatures shown in Figure 17 are indicating the presence of chiral recognition in all the polymer mixtures as well as in the monomer mixtures. Figure 18 displays the deviation ( $\Delta T$ ) of the  $S_A$ -I transition temperatures collected during the first cooling scans from the theoretical values *versus* mixture compositions. The data obtained from the monomer mixtures are also plotted in this figure. The reason for the fluctuation of data points which can be seen in this figure is not clear. Regardless of the fluctuation of the data points, the following trends can be easily observed from Figure 18. With increasing the molecular weight i.e., on going from the monomer mixtures to the polymer mixtures I, II, and III,  $\Delta T$  increases in the same order to reach a positive deviation of  $1.3^{\circ}\text{C}$  from the theoretical ideal value. The



polymer mixtures IV (polymer mixtures with the largest molecular weight) show, on the contrary, a smaller  $\Delta T$  deviation than that of the monomer mixtures. These trends clearly suggests that there is an optimum molecular weight which favors the largest chiral molecular recognition. To a certain amount of molecular weight, the polymer backbone "effect" enhances the chiral molecular recognition effect between the two enantiomeric side chain structural units through a cooperative effect.<sup>20,21</sup> However, when the molecular weight of the polymer exceeds a critical value, the polymer backbone decreases the heterochiral interaction between the enantiomeric side groups, resulting in the cancellation of chiral molecular recognition. The trend of the influence of molecular weight on the  $\Delta T$  deviation observed here suggests that the equilibrium constant of the heterochiral recognition process is molecular weight dependent. The dependence of the  $\Delta T$  deviation observed in this case resembles that of the trend observed for the dependence of the equilibrium constant of interpolymeric electron donor-acceptor complexes on molecular weight.<sup>21b</sup>

With respect to the  $S_C^*-S_A$  and  $S_X-S_C^*$  transitions of the polymer mixtures, it seems that the dependence of transition temperatures on the composition is not very clear. In Figures 11 and 14a the  $S_C^*-S_A$  transitions show a downward curvature, while in Figures 12 and 13 they look insensitive to the composition. This is probably due to the fact that the  $S_C^*-S_A$  transitions are in close proximity of the  $S_A-I$  transitions and they are partially overlapped. This can cause the inaccurate determination of the temperatures of the  $S_C^*-S_A$  transitions whose peak are much smaller than those of the  $S_A-I$  transitions.

### Copolymerization of (R)-8 and (S)-8

Copolymerization of (R)-8 with (S)-8 was performed to cover the entire range of composition. Attempts were made to synthesize poly[[(R)-8]-co-[(S)-8]] with DP=16. Since the transition temperatures of polymers are strongly dependent on their molecular weights, it is essential to synthesize polymers having identical molecular weights in order to compare their transition temperatures. This can be achieved only by a living polymerization. The copolymerization results are listed in Table IX. The yields reported in Table IX are lower

than quantitative due to the polymer losses during the purification process. However, all conversions were quantitative and therefore, the copolymer composition is identical to that of the monomer feed.<sup>10</sup> The average degrees of polymerization determined by GPC are 10-11.

The thermal transition temperatures determined by DSC are listed in Table IX and are plotted against copolymer composition in Figure 19. The copolymers poly[[(R)-8]-co-[(S)-8]] exhibit the same phase behavior as their parent homopolymers over the entire range of compositions. Although the molecular weights of these copolymers are rather well controlled, the fluctuation of the experimental data points is large and therefore, no clear trends are observed for phase transitions.

### Conclusions

In the previous publication of this series, we investigated the heterochiral molecular recognition in molecular and macromolecular pairs of diastereomeric liquid crystals based on (2R, 3S)- and (2S, 3S)-2-fluoro-3-methylpentyl 4'-(11-vinyloxyundecanyloxy)biphenyl-4-carboxylate (9) diastereomers. It was observed that chiral recognition was present in the  $S_A$  phase of monomer mixtures, however, it was canceled in polymer mixtures and in copolymers.<sup>12</sup> The current study provides the first example of heterochiral molecular recognition in enantiomeric pairs of low molar mass and side chain liquid crystalline polymers. It also presents the first experiments that demonstrate a dependence of heterochiral molecular recognition on polymer molecular weight. In the present case as in the previously reported one,<sup>12</sup> heterochiral molecular recognition was observed only in the untilted  $S_A$  phase. These results are in agreement with all other data available in the literature on low molar mass liquid crystals.<sup>1a,5,7,8</sup> On the other hand, no clear relationship between transition temperatures on composition could be obtained for copolymers. This is mostly due to the very low deviation of the transition temperature from ideal values observed for this system. This requires experiments with a more accurate polymerization technique or a pair of enantiomers which provides a larger deviation of the transition temperature as a consequence of the heterochiral recognition process.

## Acknowledgments

Financial support by the Office of Naval Research and Asahi Chemical Industry Co., Ltd., Japan is gratefully acknowledged.

## References and Notes

- (1) (a) Jacques, J.; Collet, A.; Wilen, S. H. *Enantiomers, Racemates and Resolutions*; Krieger Publishing Co.: Malabar, 1991. (b) Pirkle, W. H.; Pochapsky, T. C. *Chem. Rev.* **1989**, *89*, 347. (c) Walba, D. M.; Slater, S. C.; Thurmes, W. N.; Clark, N. A.; Handschy, M. A.; Supon, F. *J. Am. Chem. Soc.* **1986**, *108*, 5210.
- (2) (a) Arnett, E. M.; Harvey, N. G.; Rose, P. L. *Acc. Chem. Res.* **1989**, *22*, 131. (b) Rose, P. L.; Harvey, N. G.; Arnett, E. M. In *Advances in Physical Organic Chemistry* Vol. 28; Bethell, D., Ed.; Academic Press: New York, 1993; p 45. (c) Arnett, E. M.; Thompson, O. *J. Am. Chem. Soc.* **1981**, *103*, 968. (d) Harvey, N. G.; Rose, P. L.; Mirajovsky, D.; Arnett, E. M. *J. Am. Chem. Soc.* **1990**, *112*, 3547. (e) Arnett, E. M.; Gold, J. M. *J. Am. Chem. Soc.* **1982**, *104*, 636.
- (3) (a) Harvey, N.; Rose, P.; Porter, N. A.; Huff, J. B.; Arnett, E. M. *J. Am. Chem. Soc.* **1988**, *110*, 4395. (b) Heath, J. G.; Arnett, E. M. *J. Am. Chem. Soc.* **1992**, *114*, 4500.
- (4) For representative reviews on SC\* liquid crystals see: (a) Goodby, J. W. *Science* **1986**, *231*, 350. (b) Goodby, J. W. *J. Mater. Chem.* **1991**, *1*, 307. (c) Goodby, J. W., Ed. *Ferroelectric Liquid Crystals. Principles, Properties and Applications*; Gordon and Breach Science Publishers: Philadelphia, 1991. (d) LeBarny, P.; Dubois, J. C. In *Side Chain Liquid Crystal Polymers*; McArdle, C. B., Ed.; Chapman and Hall: New York, 1989; p 130. (e) Escher, C.; Wingen, R. *Adv. Mater.* **1992**, *4*, 189.
- (5) (a) Leclercq, M.; Billard, J.; Jacques, J. *Mol. Cryst. Liq. Cryst.* **1969**, *8*, 367. (b) Bahr, CH.; Heppke, G.; Sabaschus, B. *Ferroelectrics* **1988**, *84*, 103. (c) Bahr, CH.; Heppke, G.; Sabaschus, B. *Liq. Cryst.* **1991**, *9*, 31.

- (6) (a) Yamada, Y.; Mori, K.; Yamamoto, N.; Hayashi, H.; Nakamura, K.; Yamawaki, M.; Orihara, H.; Ishibashi, Y. *Jpn. J. Appl. Phys.* **1989**, *28*, L1606. (b) Takezoe, H.; Lee, J.; Chandani, A. D. L.; Gorecka, E.; Ouchi, Y.; Fukuda, A.; Terashima, K.; Furukawa, K. *Ferroelectrics* **1991**, *114*, 187. (c) Takezoe, H.; Fukuda, A.; Ikeda, A.; Takanishi, Y.; Umemoto, T.; Watanabe, J.; Iwane, H.; Hara, M.; Itoh, K. *Ferroelectrics* **1991**, *122*, 167. (d) Goodby, J. W.; Chin, E. *Liq. Cryst.* **1988**, *3*, 1245. (e) Goodby, J. W.; Patel, J. S.; Chin, E. *J. Mater. Chem.* **1992**, *2*, 197. (f) Heppke, G.; Löttsch, D.; Demus, D.; Diele, S.; Jahn, K.; Zschke, H. *Mol. Cryst. Liq. Cryst.* **1991**, *208*, 9.
- (7) (a) Goodby, J. W.; Waugh, M. A.; Stein, S. M.; Chin, E.; Pindak, R.; Patel, J. S. *J. Am. Chem. Soc.* **1989**, *111*, 8119. (b) Slaney, A. J.; Goodby, J. W. *Liq. Cryst.* **1991**, *9*, 849. (c) Goodby, J. W.; Nishiyama, I.; Slaney, A. J.; Booth, C. J.; Toyne, K. J. *Liq. Cryst.* **1993**, *14*, 37. (d) Nguyen, H. T.; Twieg, R. J.; Nabor, M. F.; Isaert, N.; Destrade, C. *Ferroelectrics* **1991**, *121*, 187. (e) Navailles, L.; Nguyen, H. T.; Barois, P.; Destrade, C.; Isaert, N. *Liq. Cryst.* **1993**, *15*, 479.
- (8) (a) Levelut, A. M.; Germain, C.; Keller, P.; Liebert, L.; Billard, J. *J. Phys., Paris* **1983**, *44*, 623. (b) Keller, P. *Mol. Cryst. Liq. Cryst. Lett.* **1984**, *102*, 295. (c) Billard, J.; Dahlgren, A.; Flatischler, K.; Lagerwall, S. T.; Otterholm, B. *J. Phys., Paris* **1985**, *46*, 1241. (d) Heppke, G.; Kleineberg, P.; Löttsch, D. *Liq. Cryst.* **1993**, *14*, 67.
- (9) Lien, S. C.; Huang, C. C.; Goodby, J. W. *Phys. Rev. A* **1984**, *29*, 1371
- (10) For a brief review on the molecular engineering of side chain LCP by living cationic polymerization see: Percec, V.; Tomazos, D. *Adv. Mater.*, **1992**, *4*, 548.
- (11) (a) Percec, V.; Zheng, Q.; Lee, M. *J. Mater. Chem.* **1991**, *1*, 611. (b) Percec, V.; Zheng, Q.; Lee, M. *J. Mater. Chem.* **1991**, *1*, 1015. (c) Percec, V.; Zheng, Q. *J. Mater. Chem.* **1992**, *2*, 475. (d) Percec, V.; Zheng, Q. *J. Mater. Chem.* **1992**, *2*, 1041.

- (12) Percec, V.; Oda, H.; Rinaldi, P. L.; Hensley, D. R. *Macromolecules*, in press.
- (13) Fu, S. J.; Birnbaum, S. M.; Greenstein, J. P. *J. Am. Chem. Soc.* **1954**, *76*, 6054.
- (14) Booth, C. J.; Goodby, J. W.; Hardy, J. P.; Lettington, O. C.; Toyne, K. J. *J. Mater. Chem.* **1993**, *3*, 821.
- (15) (a) Percec, V.; Lee, M.; Jonsson, H. *J. Polym. Sci: Part A: Polym. Chem.* **1991**, *29*, 327. (b) Percec, V.; Lee, M. *Macromolecules* **1991**, *24*, 1017. (c) Percec, V.; Lee, M.; Rinaldi, P.; Litman, V. E. *J. Polym. Sci: Part A: Polym. Chem.* **1992**, *30*, 1213.
- (16) (a) Cho, C. G.; Feit, B. A.; Webster, O. W. *Macromolecules* **1990**, *23*, 1918. (b) Cho, C. G.; Feit, B. A.; Webster, O. W. *Macromolecules* **1992**, *25*, 2081. (c) Lin, C. H.; Matyjaszewski, K. *Polym. Prepr., Am. Chem. Soc. Div. Polym. Chem.* **1990**, *31*, 599.
- (17) (a) Percec, V.; Lee, M. *Polymer* **1991**, *32*, 2862. (b) Percec, V.; Lee, M. *Polymer Bull.* **1991**, *25*, 131. (c) Percec, V.; Lee, M. *Macromolecules* **1991**, *24*, 4963. (d) Percec, V.; Lee, M. *J. Mater. Chem.* **1991**, *1*, 1007. (e) Percec, V.; Lee, M. *J. Mater. Chem.* **1992**, *2*, 617.
- (18) (a) Van Hecke, G. R. *J. Phys. Chem.* **1979**, *83*, 2344. (b) Achard, M. F.; Mauzac, M.; Richard, H.; Sigaud, G.; Hardouin, F. *Eur. Polym. J.* **1989**, *25*, 593.
- (19) (a) Percec, V.; Lee, M. *J. Mater. Chem.* **1991**, *1*, 1007. (b) Percec, V.; Lee, M.; Zheng, Q. *Liq. Cryst.* **1992**, *12*, 715. (c) Percec, V.; Johansson, G. *J. Mater. Chem.* **1993**, *3*, 83.
- (20) (a) Ciardelli, F.; Salvadori, P. *Pure Appl. Chem.* **1985**, *57*, 931. (b) Selegny, E., Ed. *Optically Active Polymers*; D. Reidel Publishing Co.: Dordrecht, The Netherlands, 1979. (c) Matsuzaki, K.; Watanabe, T. *Makromol. Chem.* **1971**, *146*, 109.
- (21) (a) Pugh, C.; Rodriguez-Parada, J.; Percec, V. *J. Polym. Sci: Part A: Polym. Chem.*, **1986**, *24*, 747; (b) V. Percec, H. G. Schild, J. M. Rodriguez-Parada and C. Pugh. *J. Polym. Sci: Part A: Polym. Chem.* **1988**, *26*, 935.

### Scheme and Figure Captions

Scheme 1. Synthesis of monomers **(R)-8** and **(S)-8**.

Scheme 2. Cationic polymerization of **(R)-8** and **(S)-8**.

Scheme 3. Cationic copolymerization of **(S)-8** with **(2S,3S)-9**.

Figure 1.  $^1\text{H}$ -NMR spectra of monomer **(R)-8** (a) and polymer poly[**(R)-8**] (DP=5.7) (b).

Figure 2.  $^1\text{H}$ -NMR spectra of monomer mixture and monomers with  $\text{Eu}(\text{hfc})_3$ :

(a) **(R)-8**/**(S)-8** (50/50 mixture); (b) **(R)-8**; (c) **(S)-8**.

Figure 3. Dependence of the number-average molecular weight ( $M_n$ ) and polydispersity ( $M_w/M_n$ ) of poly[**(R)-8**] (open symbols) and poly[**(S)-8**] (closed symbols) determined by GPC on the  $[\text{M}]_0/[\text{I}]_0$  ratio.

Figure 4. DSC thermograms (20°C/min) of poly[**(R)-8**] with different DP: (a) first heating scans; (b) second heating scans; (c) first cooling scans.

Figure 5. DSC thermograms (20°C/min) of poly[**(S)-8**] with different DP: (a) first heating scans; (b) second heating scans; (c) first cooling scans.

Figure 6. Dependence of phase transition temperatures on the degree of polymerization of poly[**(R)-8**] (open) and poly[**(S)-8**] (closed): (a) data from the first heating scans; (b) data from the second heating scans; (c) data from the first cooling scans.

Figure 7. Representative optical polarized micrographs of the  $S_A$  mesophase displayed by poly[**(S)-8**] (DP=11.8) upon heating to 78°C (100x).

Figure 8. DSC thermograms (20°C/min) of poly[[**(S)-8**]-*co*-[(**2S,3S**)-**9**]]X/Y with different compositions: (a) first heating scans; (b) second heating scans; (c) first cooling scans.

Figure 9. Dependence of phase transition temperatures on the composition of poly[[**(S)-8**]-*co*-[(**2S,3S**)-**9**]]

- Figure 10. Dependence of phase transition temperatures on the composition of the binary mixtures of (R)-8 with (S)-8: (a) data from the first heating scans; (b) data from the second heating scans; (c) data from the first cooling scans.
- Figure 11. Dependence of phase transition temperatures on the composition of polymer mixture I [poly[(R)-8] (DP=5.7) and poly[(S)-8] (DP=6.1)]: (a) data from the first heating scans; (b) data from the second heating scans; (c) data from the first cooling scans.
- Figure 12. The dependence of phase transition temperatures on the composition of polymer mixture II [poly[(R)-8] (DP=6.7) and poly[(S)-8] (DP=6.9)]: (a) data from the first heating scans; (b) data from the second heating scans; (c) data from the first cooling scans.
- Figure 13. The dependence of phase transition temperatures on the composition of polymer mixture III [poly[(R)-8] (DP=9.6) and poly[(S)-8] (DP=10.1)]: (a) data from the first heating scans; (b) data from the second heating scans; (c) data from the first cooling scans.
- Figure 14. The dependence of phase transition temperatures on the composition of polymer mixture IV [poly[(R)-8] (DP=15.1) with poly[(S)-8] (DP=14.3)]: (a) data from the first heating scans; (b) data from the second heating scans; (c) data from the first cooling scans.
- Figure 15. DSC thermograms (10°C/min) of the binary mixtures of monomer (R)-8 (X) with monomer (S)-8 (Y): (a) first heating scans; (b) second heating scans; (c) first cooling scans.
- Figure 16. DSC thermograms (10°C/min) of polymer mixture III [poly[(R)-8] (DP=9.6) and poly[(S)-8] (DP=10.1)]: (a) first heating scans; (b) second heating scans; (c) first cooling scans.
- Figure 17. Dependence of the  $S_A$ -I transition temperatures from experimental data (open symbols) and from the Schröder-van Laar equation (closed symbols) on the

composition of the binary mixtures of poly[(R)-8] with poly[(S)-8]: (a) data from the first heating scans; b) data from the second heating scans; (c) data from the first cooling scans.

Figure 18. Deviation of the  $S_A$ -I transition temperatures collected during the first cooling scans from the theoretical ideal values ( $\Delta T$ ) *versus* composition.

Figure 19. Dependence of the phase transition temperatures on the composition of poly[(R)-8-co-(S)-8] (DP=10.1-10.7): (a) data from the first heating scans; (b) data from the second heating scans; (c) data from the first cooling scans.



Table 1. Cationic Polymerization of (R)-2-Chloro-4-Methylpentyl 4'-(8-Vinyloxyoctyloxy)bi(phenyl)-4-Carboxylate ((R)-8) and Characterization of the Resulting Polymers<sup>a</sup>

Sample No.	[M] <sub>0</sub> /[I] <sub>0</sub>	Polymer Yield(%)	M <sub>n</sub> × 10 <sup>-3</sup>	M <sub>w</sub> /M <sub>n</sub>	DP	Phase transitions (°C) and corresponding enthalpy changes (kcal/mru) <sup>b</sup>	
						heating	cooling
1	3	52.1	1.99	1.12	4.1	S <sub>C</sub> * 53.1(0.34) S <sub>A</sub> 56.0(1.03) I S <sub>C</sub> * 53.0(0.32) S <sub>A</sub> 56.3(1.00) I	1 47.3 (-1.21) S <sub>C</sub> *
2	5	78.6	2.76	1.14	5.7	S <sub>C</sub> * 59.3(0.31) S <sub>A</sub> 62.2(0.84) I S <sub>C</sub> * 59.0(0.32) S <sub>A</sub> 62.1(0.80) I	1 52.4 (-1.04) S <sub>C</sub> *
3	8	68.9	3.40	1.23	7.0	S <sub>C</sub> * 64.6(0.27) S <sub>A</sub> 69.4(0.82) I S <sub>C</sub> * 64.2(0.27) S <sub>A</sub> 70.5(0.81) I	1 60.4 (-0.97) S <sub>C</sub> *
4	12	91.0	4.25	1.22	8.7	K 41.5(0.58) S <sub>C</sub> * 72.1(0.27) S <sub>A</sub> 77.1(0.92) I S <sub>X</sub> 35.1(0.08) S <sub>C</sub> * 71.1(0.29) S <sub>A</sub> 76.2(0.82) I	1 66.1 (-1.12) S <sub>C</sub> * 26.2 (-0.07) S <sub>X</sub>
5	16	80.3	4.75	1.16	9.6	S <sub>X</sub> 43.2(0.07) S <sub>C</sub> * 74.5(0.33) S <sub>A</sub> 78.8(0.71) I S <sub>X</sub> 41.5(0.08) S <sub>C</sub> * 74.2(0.28) S <sub>A</sub> 78.2(0.72) I	1 68.1 (-1.04) S <sub>C</sub> * 33.6 (-0.08) S <sub>X</sub>
6	20	78.3	5.93	1.16	12.2	S <sub>X</sub> 52.1(0.11) S <sub>C</sub> * 77.3(0.30) S <sub>A</sub> 81.7(0.78) I S <sub>X</sub> 50.8(0.12) S <sub>C</sub> * 76.9(0.28) S <sub>A</sub> 81.1(0.77) I	1 71.4 (-1.06) S <sub>C</sub> * 43.5 (-0.10) S <sub>X</sub>
7	30	82.5	7.33	1.20	15.1	S <sub>X</sub> 62.9(0.16) S <sub>C</sub> * 81.3(0.28) S <sub>A</sub> 85.9(0.82) I S <sub>X</sub> 61.7(0.18) S <sub>C</sub> * 80.5(0.30) S <sub>A</sub> 84.8(0.75) I	1 75.0 (-1.03) S <sub>C</sub> * 54.3 (-0.16) S <sub>X</sub>

<sup>a</sup> Polymerization temperature: 0°C; polymerization solvent: methylene chloride; [M]<sub>0</sub>=0.224; [M]<sub>2</sub>S<sub>1</sub>/[I]<sub>0</sub>=10; polymerization time: 1 hour.

<sup>b</sup> Data on the first line are from first heating and cooling scans. Data on the second line are from second heating scan. Heating and cooling rates are 20°C/min.

Table II Cationic Polymerization of (S)-2-Chloro-4-Methylpentyl-4'-(8-Vinyloxyoctyloxy)biphenyl-4-Carboxylate ((S)-8) and Characterization of the Resulting Polymers<sup>a</sup>

Sample No.	[M] <sub>0</sub> /[I] <sub>0</sub>	Polymer Yield(%)	Max(°C)	Mw/Mn	DP	Phase transitions (°C) and corresponding enthalpy changes (kcal/mole) <sup>b</sup>	
						heating	cooling
1	3	47.1	1.91	1.12	3.9	S <sub>C</sub> * 52.3(0.31) S <sub>A</sub> 56.0(0.90) I	147.7(1.06) S <sub>C</sub> *
						S <sub>C</sub> * 52.1(0.27) S <sub>A</sub> 56.1(0.89) I	
2	5	75.7	2.32	1.23	4.8	S <sub>C</sub> * 56.7(1°) S <sub>A</sub> 59.5(1.13) I	149.6(1.02) S <sub>C</sub> *
						S <sub>C</sub> * 55.8(0.32) S <sub>A</sub> 59.2(0.77) I	
3	8	82.8	2.97	1.23	6.1	S <sub>C</sub> * 63.0(0.30) S <sub>A</sub> 66.9(0.77) I	157.4(1.01) S <sub>C</sub> *
						S <sub>C</sub> * 62.6(0.27) S <sub>A</sub> 67.0(0.80) I	
4	12	83.2	3.38	1.21	6.9	K 42.0(0.16) S <sub>C</sub> * 68.1(0.34) S <sub>A</sub> 72.0(0.98) I	161.4(1.21) S <sub>C</sub> *
						S <sub>X</sub> 18.6(0.08) S <sub>C</sub> * 67.1(0.34) S <sub>A</sub> 71.3(0.92) I	
5	12	89.8	4.12	1.19	8.5	K 42.4(0.62) S <sub>C</sub> * 72.0(0.29) S <sub>A</sub> 77.0(0.90) I	166.1 (-1.05) S <sub>C</sub> * 26.1 (-0.07) S <sub>X</sub>
						S <sub>X</sub> 34.2(0.07) S <sub>C</sub> * 71.5(0.28) S <sub>A</sub> 76.2(0.80) I	
6	20	70.8	5.77	1.16	11.8	S <sub>X</sub> 51.1(0.13) S <sub>C</sub> * 76.9(0.37) S <sub>A</sub> 81.3(0.72) I	170.9 (-1.03) S <sub>C</sub> * 41.5 (-0.10) S <sub>X</sub>
						S <sub>X</sub> 49.2(0.13) S <sub>C</sub> * 76.1(0.34) S <sub>A</sub> 80.3(0.73) I	
7	30	81.1	7.20	1.20	14.8	S <sub>X</sub> 60.6(0.17) S <sub>C</sub> * 80.0(0.31) S <sub>A</sub> 84.5(0.78) I	173.4 (-1.06) S <sub>C</sub> * 51.2 (-0.14) S <sub>X</sub>
						S <sub>X</sub> 58.6(0.16) S <sub>C</sub> * 78.9(0.28) S <sub>A</sub> 83.1(0.77) I	

<sup>a</sup> Polymerization temperature: 0°C; polymerization solvent: methylene chloride; [M]<sub>0</sub>=0.224; [Me<sub>2</sub>Si]/[I]<sub>0</sub>=10; polymerization time: 1 hour.

<sup>b</sup> Data on the first line are from first heating and cooling scans. Data on the second line are from second heating scan. Heating and cooling rates are 20°C/min.

<sup>c</sup> Overlapped peak

Table III. Cationic Copolymerization of (S)-8 with (2S,3S)-9 and Characterization of the Resulting Polymers<sup>a</sup>

[(S)-8]/ [(2S,3S)-9] (mol/mol)	Polymer Yield(%)	Mn $\times 10^{-3}$	Mw/Mn	DP	Phase transitions (°C) and corresponding enthalpy changes (kcal/mole) <sup>b</sup>	
					heating	cooling
0/100	74.0	5.78	1.10	11.3	K 53.0(+) <sup>c</sup> S <sub>X</sub> 57.7(1.39) S <sub>C</sub> * 92.4(0.04) S <sub>A</sub> 120.9(0.17) I	I 111.7(-1.17) S <sub>A</sub> 85.6(0.04) S <sub>C</sub> * 48.2(-0.88) S <sub>X</sub>
10/90	79.9	5.37	1.23	10.5	S <sub>X</sub> 57.2(1.02) S <sub>C</sub> * 92.1(0.03) S <sub>A</sub> 120.3(0.17) I K 50.4(1.59) S <sub>C</sub> * 81.4(0.03) S <sub>A</sub> 116.5(0.31) I	I 107.3(-1.28) S <sub>A</sub> 74.6(0.04) S <sub>C</sub> * 39.8(-0.91) S <sub>X</sub>
25/75	73.8	5.63	1.17	11.1	S <sub>X</sub> 49.2(1.06) S <sub>C</sub> * 81.7(0.04) S <sub>A</sub> 116.4(1.35) I K 42.1(1.46) S <sub>C</sub> * 70.8(0.04) S <sub>A</sub> 111.0(1.28) I	I 101.4(-1.21) S <sub>A</sub> 63.7(0.03) S <sub>C</sub> * 25.7(-0.71) S <sub>X</sub>
40/60	67.9	5.57	1.21	11.1	S <sub>X</sub> 34.8(0.86) S <sub>C</sub> * 70.6(0.04) S <sub>A</sub> 110.5(1.26) I K 38.5(1.27) S <sub>C</sub> * 65.9(0.04) S <sub>A</sub> 106.5(1.14) I	I 96.4(-1.10) S <sub>A</sub> 58.0(0.05) S <sub>C</sub> * 21.7(-0.67) S <sub>X</sub>
60/40	54.1	5.94	1.17	11.9	S <sub>X</sub> 31.2(0.79) S <sub>C</sub> * 65.1(0.05) S <sub>A</sub> 106.3(1.13) I S <sub>X</sub> 21.5(+) <sup>c</sup> K 38.9(0.58) S <sub>C</sub> * 63.9(0.11) S <sub>A</sub> 94.2(0.94) I	I 84.2(-0.92) S <sub>A</sub> 57.0(0.10) S <sub>C</sub> * 11.3(-0.13) S <sub>X</sub>
75/25	81.4	5.94	1.21	12.0	S <sub>X</sub> 21.5(0.52) S <sub>C</sub> * 63.4(0.10) S <sub>A</sub> 93.7(0.95) I S <sub>X</sub> 33.6(0.07) S <sub>C</sub> * 69.4(0.15) S <sub>A</sub> 89.7(0.81) I	I 79.7(-0.79) S <sub>A</sub> 62.1(-0.14) S <sub>C</sub> * 22.4(-0.06) S <sub>X</sub>
90/10	69.8	5.95	1.18	12.1	S <sub>X</sub> 31.3(0.06) S <sub>C</sub> * 68.7(0.17) S <sub>A</sub> 89.2(0.83) I S <sub>X</sub> 34.9(0.06) S <sub>C</sub> * 70.1(0.18) S <sub>A</sub> 81.8(0.73) I	I 72.1(-0.69) S <sub>A</sub> 63.2(-0.17) S <sub>C</sub> * 23.9(-0.05) S <sub>X</sub>
100/0	70.8	5.77	1.16	11.8	S <sub>X</sub> 33.1(0.06) S <sub>C</sub> * 69.8(0.20) S <sub>A</sub> 81.7(0.72) I S <sub>X</sub> 51.1(0.13) S <sub>C</sub> * 76.9(0.37) S <sub>A</sub> 81.3(0.72) I	I 70.9(-1.03) S <sub>C</sub> * 41.5(-0.10) S <sub>X</sub>
					S <sub>X</sub> 49.2(0.13) S <sub>C</sub> * 76.1(0.34) S <sub>A</sub> 80.3(0.73) I	

<sup>a</sup> Polymerization temperature: 0°C; polymerization solvent: methylene chloride; [M]<sub>0</sub>=[(S)-8]+[(2S,3S)-9]=0.224;[N]<sub>0</sub>/[I]<sub>0</sub>=20; [Me<sub>2</sub>Si]<sub>0</sub>/[I]<sub>0</sub>=10; polymerization time: 1 hour.<sup>b</sup> Heating and cooling rates are 20°C/min.<sup>c</sup> Data on the first line are from first heating and cooling scans. Data on the second line are from second heating scan.

\* Overlapped peak.

Table IV. Characterization of the Binary Mixtures of Monomers (R)-8 with (S)-8

(R)-8/(S)-8 (mol)/(mol)	Phase transitions (°C) and corresponding enthalpy changes (kcal/mol) <sup>a</sup>	
	heating	cooling
0/100	K 52.8 (11.80) I K 38.6 (3.10) K 40.1 (-1.50) K 49.5 (8.06) I	I 29.7 (-1.05) S <sub>A</sub> 17.4 (-0.05) S <sub>C</sub> * 7.9 (-5.52) K
19.8/80.2	K 48.0 (11.40) I K 45.4 (10.73) I	I 30.2 (-1.03) S <sub>A</sub> 17.5 (-0.05) S <sub>C</sub> * -4.8 (-5.60) K
39.0/61.0	K 49.4 (11.27) I K 48.3 (10.79) I	I 30.2 (-1.03) S <sub>A</sub> 17.3 (-0.06) S <sub>C</sub> * -6.4 (-4.87) K
44.8/55.2	K 50.1 (11.49) I K 48.8 (11.06) I	I 30.3 (-1.05) S <sub>A</sub> 17.2 (-0.06) S <sub>C</sub> * -6.6 (-4.99) K
49.9/50.1	K 50.2 (12.05) I K 42.5 (0.37) K 48.4 (10.93) I	I 30.2 (-1.10) S <sub>A</sub> 16.9 (-0.07) S <sub>C</sub> * -6.6 (-5.35) K
55.0/45.0	K 49.5 (11.78) I K 48.6 (11.41) I	I 30.2 (-1.07) S <sub>A</sub> 16.9 (-0.07) S <sub>C</sub> * -6.7 (-5.06) K
59.4/40.6	K 49.4 (11.22) I K 48.5 (11.12) I	I 30.2 (-1.06) S <sub>A</sub> 16.9 (-0.07) S <sub>C</sub> * -6.7 (-5.13) K
80.0/20.0	K 48.2 (11.45) I K 45.8 (10.93) I	I 30.1 (-1.06) S <sub>A</sub> 17.3 (-0.05) S <sub>C</sub> * -5.3 (-5.67) K
100/0	K 51.4 (-11.76) I K 38.6 (3.96) K 39.9 (-3.78) K 49.0 (9.14) I	I 29.9 (-1.04) S <sub>A</sub> 17.4 (-0.06) S <sub>C</sub> * 6.7 (-5.86) K

<sup>a</sup> Data on the first line are from first heating and cooling scans. Data on the second line are from second heating scan. Heating and cooling rates are 10°C/min.

Table V. Characterization of the Binary Mixtures of Poly[(R)-8] (DP=5.7,  $M_w/M_n=1.14$ ) with Poly[(S)-8] (DP=6.1,  $M_w/M_n=1.23$ ) (Polymer Mixture I)

poly[(R)-8]/poly[(S)-8] (mol)/(mol)	Phase transitions (°C) and corresponding enthalpy changes (kcal/mru) <sup>a</sup>	
	heating	cooling
0/100	S <sub>C</sub> * 59.6 (0.19) S <sub>A</sub> 64.1 (0.75) I S <sub>C</sub> * 59.2 (0.27) S <sub>A</sub> 64.1 (0.77) I	I 57.6 (-1.02) S <sub>A</sub> 56.3 (†b) S <sub>C</sub> *
10.8/89.2	S <sub>C</sub> * 58.7 (0.19) S <sub>A</sub> 63.3 (0.73) I S <sub>C</sub> * 58.7 (0.25) S <sub>A</sub> 63.8 (0.78) I	I 57.4 (-0.98) S <sub>A</sub> 55.4 (†b) S <sub>C</sub> *
20.0/80.0	S <sub>C</sub> * 58.1 (0.20) S <sub>A</sub> 62.6 (0.79) I S <sub>C</sub> * 58.0 (0.26) S <sub>A</sub> 63.1 (0.83) I	I 57.0 (-1.06) S <sub>A</sub> 54.8 (†b) S <sub>C</sub> *
50.2/49.8	S <sub>C</sub> * 56.8 (0.21) S <sub>A</sub> 61.2 (0.71) I S <sub>C</sub> * 56.4 (0.29) S <sub>A</sub> 61.4 (0.80) I	I 55.3 (-1.03) S <sub>C</sub> *
59.3/40.7	S <sub>C</sub> * 56.8 (0.20) S <sub>A</sub> 61.2 (0.81) I S <sub>C</sub> * 56.7 (0.28) S <sub>A</sub> 61.8 (0.82) I	I 55.4 (-1.07) S <sub>A</sub> 53.8 (†b) S <sub>C</sub> *
80.7/19.3	S <sub>C</sub> * 56.0 (0.19) S <sub>A</sub> 60.0 (0.83) I S <sub>C</sub> * 56.0 (0.29) S <sub>A</sub> 60.6 (0.94) I	I 54.3 (-1.17) S <sub>C</sub> *
100/0	S <sub>C</sub> * 55.6 (0.20) S <sub>A</sub> 59.1 (0.80) I S <sub>C</sub> * 55.4 (0.30) S <sub>A</sub> 59.2 (0.85) I	I 52.9 (-1.06) S <sub>C</sub> *

<sup>a</sup> Data on the first line are from first heating and cooling scans. Data on the second line are from second heating scan.

Heating and cooling rates are 10°C/min.

<sup>b</sup> Overlapped peak.

Table VI. Characterization of the Binary Mixtures of Poly[(R)-8] (DP=6.7,  $M_w/M_n=1.23$ ) with Poly[(S)-8] (DP=6.9,  $M_w/M_n=1.21$ ) (Polymer Mixture II).

poly[(R)-8]/poly[(S)-8] (mol)/(mol)	Phase transitions (°C) and corresponding enthalpy changes (kcal/mol) <sup>a</sup>	
	heating	cooling
0/100	K 41.6 (0.38) S <sub>C</sub> * 63.9 (0.25) S <sub>A</sub> 68.8 (0.78) I S <sub>X</sub> 15.5 (0.05) S <sub>C</sub> * 63.6 (0.25) S <sub>A</sub> 68.4 (0.74) I	I 61.8 (-0.98) S <sub>C</sub> *
21.4/78.6	S <sub>X</sub> 14.0 (0.06) S <sub>C</sub> * 62.7 (0.25) S <sub>A</sub> 67.2 (0.76) I S <sub>X</sub> 12.4 (0.03) S <sub>C</sub> * 62.5 (0.24) S <sub>A</sub> 67.7 (0.75) I	I 61.3 (-0.99) S <sub>A</sub> 59.1 († <sup>b</sup> ) S <sub>C</sub> *
39.1/60.9	S <sub>C</sub> * 62.0 (0.23) S <sub>A</sub> 66.9 (0.76) I S <sub>X</sub> 11.2 (0.02) S <sub>C</sub> * 61.7 (0.25) S <sub>A</sub> 67.3 (0.76) I	I 61.1 (-1.00) S <sub>A</sub> 58.5 († <sup>b</sup> ) S <sub>C</sub> *
45.6/54.4	S <sub>X</sub> 11.2 (0.01) S <sub>C</sub> * 61.6 (0.25) S <sub>A</sub> 66.6 (0.76) I S <sub>X</sub> 11.5 (0.03) S <sub>C</sub> * 61.5 (0.29) S <sub>A</sub> 67.2 (0.78) I	I 61.2 (-1.00) S <sub>A</sub> 58.3 († <sup>b</sup> ) S <sub>C</sub> *
50.1/49.9	S <sub>C</sub> * 61.5 (0.28) S <sub>A</sub> 66.4 (0.80) I S <sub>X</sub> 10.2 (0.02) S <sub>C</sub> * 61.5 (0.22) S <sub>A</sub> 67.1 (0.76) I	I 61.2 (-0.95) S <sub>A</sub> 58.2 († <sup>b</sup> ) S <sub>C</sub> *
54.9/45.1	S <sub>X</sub> 10.1 (0.04) S <sub>C</sub> * 61.5 (0.24) S <sub>A</sub> 66.2 (0.75) I S <sub>X</sub> 10.2 (0.03) S <sub>C</sub> * 61.3 (0.25) S <sub>A</sub> 67.1 (0.77) I	I 60.8 (-0.97) S <sub>A</sub> 57.9 († <sup>b</sup> ) S <sub>C</sub> *
59.0/41.0	S <sub>X</sub> 11.1 (0.02) S <sub>C</sub> * 61.8 (0.25) S <sub>A</sub> 66.7 (0.80) I S <sub>X</sub> 10.7 (0.02) S <sub>C</sub> * 61.6 (0.23) S <sub>A</sub> 67.2 (0.76) I	I 61.1 (-0.95) S <sub>A</sub> 58.2 († <sup>b</sup> ) S <sub>C</sub> *
88.4/11.6	S <sub>C</sub> * 60.1 (0.24) S <sub>A</sub> 64.9 (0.78) I S <sub>C</sub> * 59.9 (0.26) S <sub>A</sub> 65.4 (0.78) I	I 59.6 (-1.00) S <sub>A</sub> 56.9 († <sup>b</sup> ) S <sub>C</sub> *
100/0	S <sub>C</sub> * 59.6 (0.22) S <sub>A</sub> 64.2 (0.74) I S <sub>C</sub> * 59.5 (0.26) S <sub>A</sub> 64.8 (0.77) I	I 58.8 (-0.98) S <sub>A</sub> 56.4 († <sup>b</sup> ) S <sub>C</sub> *

<sup>a</sup> Data on the first line are from first heating and cooling scans. Data on the second line are from second heating scan.

Heating and cooling rates are 10°C/min.

<sup>b</sup> Overlapped peak.

Table VII. Characterization of the Binary Mixtures of Poly[(R)-8] (DP=9.6,  $M_w/M_n=1.16$ ) with Poly[(S)-8] (DP=10.1,  $M_w/M_n=1.16$ ) (Polymer Mixture III)

poly[(R)-8]/poly[(S)-8] (mol)/(mol)	Phase transitions (°C) and corresponding enthalpy changes (kcal/mol) <sup>a</sup>	
	heating	cooling
0/100	S <sub>X</sub> 42.2(0.21) K 45.5(† <sup>b</sup> ) S <sub>C</sub> * 71.1(0.17) S <sub>A</sub> 75.3(0.80) I S <sub>X</sub> 40.6(0.08) S <sub>C</sub> * 70.7(0.28) S <sub>A</sub> 75.2(0.71) I	I 69.0(-1.00) S <sub>A</sub> 67.3(† <sup>b</sup> ) S <sub>C</sub> * 35.9(-0.07) S <sub>X</sub>
27.1/72.9	S <sub>X</sub> 40.1(0.05) K 44.8(0.02) S <sub>C</sub> * 70.7(0.21) S <sub>A</sub> 75.0(0.76) I S <sub>X</sub> 39.4(0.08) S <sub>C</sub> * 70.7(0.25) S <sub>A</sub> 75.4(0.78) I	I 69.7(-0.77) S <sub>A</sub> 67.5(-0.27) S <sub>C</sub> * 35.2(-0.08) S <sub>X</sub>
34.4/65.6	S <sub>X</sub> 40.2(0.06) S <sub>C</sub> * 70.7(0.21) S <sub>A</sub> 75.3(0.76) I S <sub>X</sub> 39.4(0.08) S <sub>C</sub> * 70.8(0.29) S <sub>A</sub> 75.6(0.75) I	I 69.9(-0.74) S <sub>A</sub> 67.5(-0.26) S <sub>C</sub> * 35.7(-0.08) S <sub>X</sub>
44.3/55.7	S <sub>X</sub> 39.9(0.04) S <sub>C</sub> * 70.6(0.20) S <sub>A</sub> 75.1(0.74) I S <sub>X</sub> 39.0(0.08) S <sub>C</sub> * 70.6(0.26) S <sub>A</sub> 75.5(0.75) I	I 69.7(-0.73) S <sub>A</sub> 67.4(-0.25) S <sub>C</sub> * 35.3(-0.10) S <sub>X</sub>
47.1/52.9	S <sub>X</sub> 40.0(0.06) S <sub>C</sub> * 70.8(0.23) S <sub>A</sub> 75.3(0.75) I S <sub>X</sub> 39.4(0.09) S <sub>C</sub> * 70.8(0.23) S <sub>A</sub> 75.7(0.80) I	I 70.2(-0.75) S <sub>A</sub> 67.6(-0.26) S <sub>C</sub> * 36.0(-0.07) S <sub>X</sub>
49.9/50.1	S <sub>X</sub> 39.4(0.05) S <sub>C</sub> * 70.8(0.20) S <sub>A</sub> 75.4(0.81) I S <sub>X</sub> 39.4(0.09) S <sub>C</sub> * 70.7(0.29) S <sub>A</sub> 75.6(0.77) I	I 69.7(-0.76) S <sub>A</sub> 67.3(-0.28) S <sub>C</sub> * 35.3(-0.08) S <sub>X</sub>
55.5/44.5	S <sub>X</sub> 39.8(0.05) S <sub>C</sub> * 70.9(0.19) S <sub>A</sub> 75.4(0.72) I S <sub>X</sub> 39.4(0.08) S <sub>C</sub> * 70.8(0.27) S <sub>A</sub> 75.8(0.74) I	I 69.9(-0.74) S <sub>A</sub> 67.6(-0.27) S <sub>C</sub> * 35.9(-0.08) S <sub>X</sub>
74.9/25.1	S <sub>X</sub> 39.4(† <sup>b</sup> ) K 44.3(0.10) S <sub>C</sub> * 70.8(0.23) S <sub>A</sub> 75.2(0.73) I S <sub>X</sub> 38.6(0.08) S <sub>C</sub> * 70.7(0.28) S <sub>A</sub> 75.6(0.74) I	I 69.6(-0.71) S <sub>A</sub> 67.4(-0.26) S <sub>C</sub> * 34.6(-0.06) S <sub>X</sub>
100/0	K 43.8(0.37) S <sub>C</sub> * 70.8(0.26) S <sub>A</sub> 75.0(0.73) I S <sub>X</sub> 39.0(0.08) S <sub>C</sub> * 70.7(0.29) S <sub>A</sub> 75.2(0.75) I	I 68.8(-1.00) S <sub>A</sub> 67.3(† <sup>b</sup> ) S <sub>C</sub> * 34.0(-0.07) S <sub>X</sub>

<sup>a</sup> Data on the first line are from first heating and cooling scans. Data on the second line are from second heating scan.

Heating and cooling rates are 10°C/min.

<sup>b</sup> Overlapped peak.

Table VIII. Characterization of the Binary Mixtures of Poly[(R)-8] (DP=15.1,  $M_w/M_n=1.20$ ) with Poly[(S)-8] (DP=14.3,  $M_w/M_n=1.19$ ) (Polymer Mixture IV)

poly[(R)-8]/poly[(S)-8] (mol)/(mol)	Phase transitions (°C) and corresponding enthalpy changes (kcal/mru) <sup>a</sup>			
	heating		cooling	
0/100	S <sub>X</sub> 55.4 (0.09) S <sub>C</sub> * 76.4 (0.25) S <sub>A</sub> 81.2 (0.73) I S <sub>X</sub> 53.4 (0.13) S <sub>C</sub> * 75.2 (0.32) S <sub>A</sub> 80.3 (0.70) I		I 73.8 (-0.98) S <sub>A</sub> 72.6 († <sup>b</sup> ) S <sub>C</sub> * 49.1 (-0.12) S <sub>X</sub>	
11.5/88.5	S <sub>X</sub> 54.9 (0.12) S <sub>C</sub> * 76.4 (0.29) S <sub>A</sub> 81.3 (0.78) I S <sub>X</sub> 54.2 (0.12) S <sub>C</sub> * 76.0 (0.31) S <sub>A</sub> 80.7 (0.71) I		I 74.3 (-0.74) S <sub>A</sub> 72.7 (-0.26) S <sub>C</sub> * 50.0 (-0.12) S <sub>X</sub>	
21.8/78.2	S <sub>X</sub> 55.5 (0.13) S <sub>C</sub> * 76.6 (0.26) S <sub>A</sub> 81.6 (0.75) I S <sub>X</sub> 54.9 (0.13) S <sub>C</sub> * 76.4 (0.29) S <sub>A</sub> 81.3 (0.77) I		I 74.2 (-0.98) S <sub>A</sub> 72.6 († <sup>b</sup> ) S <sub>C</sub> * 49.9 (-0.12) S <sub>X</sub>	
38.9/61.1	S <sub>X</sub> 56.5 (0.15) S <sub>C</sub> * 76.9 (0.28) S <sub>A</sub> 82.0 (0.79) I S <sub>X</sub> 55.4 (0.15) S <sub>C</sub> * 76.5 (0.29) S <sub>A</sub> 81.4 (0.72) I		I 75.1 (-0.72) S <sub>A</sub> 73.2 (-0.30) S <sub>C</sub> * 51.6 (-0.15) S <sub>X</sub>	
50.0/50.0	S <sub>X</sub> 57.1 (0.13) S <sub>C</sub> * 77.1 (0.26) S <sub>A</sub> 82.2 (0.76) I S <sub>X</sub> 55.7 (0.15) S <sub>C</sub> * 76.7 (0.24) S <sub>A</sub> 81.6 (0.72) I		I 75.3 (-0.76) S <sub>A</sub> 73.4 (-0.20) S <sub>C</sub> * 52.2 (-0.14) S <sub>X</sub>	
59.2/40.8	S <sub>X</sub> 57.5 (0.16) S <sub>C</sub> * 77.4 (0.29) S <sub>A</sub> 82.3 (0.79) I S <sub>X</sub> 56.6 (0.16) S <sub>C</sub> * 77.0 (0.28) S <sub>A</sub> 82.0 (0.74) I		I 75.9 (-0.71) S <sub>A</sub> 73.8 (-0.28) S <sub>C</sub> * 53.0 (-0.15) S <sub>X</sub>	
80.9/19.1	S <sub>X</sub> 58.8 (0.14) S <sub>C</sub> * 77.9 (0.28) S <sub>A</sub> 82.6 (0.78) I S <sub>X</sub> 58.2 (0.16) S <sub>C</sub> * 77.6 (0.31) S <sub>A</sub> 82.2 (0.71) I		I 75.8 (-0.71) S <sub>A</sub> 74.1 (-0.26) S <sub>C</sub> * 53.9 (-0.15) S <sub>X</sub>	
100/0	S <sub>X</sub> 60.4 (0.15) S <sub>C</sub> * 78.5 (0.30) S <sub>A</sub> 82.9 (0.75) I S <sub>X</sub> 59.7 (0.17) S <sub>C</sub> * 78.2 (0.33) S <sub>A</sub> 82.5 (0.71) I		I 76.4 (-0.68) S <sub>A</sub> 74.7 (-0.31) S <sub>C</sub> * 55.2 (-0.16) S <sub>X</sub>	

<sup>a</sup> Data on the first line are from first heating and cooling scans. Data on the second line are from second heating scan.  
Heating and cooling rates are 10°C/min.

<sup>b</sup> Overlapped peak.



Table IX. Cationic Copolymerization of (R)-8 with (S)-8 and Characterization of the Resulting Polymers<sup>a</sup>

[(R)-8]/[(S)-8] (mol/mol)	Polymer Yield(%)	Mn $\times 10^{-3}$	Mw/Mn	DP	Phase transitions (°C) and corresponding enthalpy changes (kcal/mru) <sup>b</sup>		
					heating	cooling	
0/100	76.2	4.99	1.16	10.1	S <sub>X</sub> 42.2(0.21) K 45.5(† <sup>c</sup> ) S <sub>C</sub> * 71.1(0.17) S <sub>A</sub> 75.3(0.80) I	I 69.0(-1.00) S <sub>A</sub> 67.3(† <sup>c</sup> ) S <sub>C</sub> * 35.9(-0.07) S <sub>X</sub>	
					S <sub>X</sub> 40.6(0.08) S <sub>C</sub> * 70.7(0.28) S <sub>A</sub> 75.2(0.71) I		
20/80	79.3	5.17	1.15	10.6	S <sub>X</sub> 44.7(0.10) S <sub>C</sub> * 72.9(0.27) S <sub>A</sub> 77.7(0.86) I	I 70.7(-1.07) S <sub>A</sub> 69.1(† <sup>c</sup> ) S <sub>C</sub> * 39.5(-0.07) S <sub>X</sub>	
					S <sub>X</sub> 44.1(0.10) S <sub>C</sub> * 72.8(0.25) S <sub>A</sub> 77.4(0.82) I		
40/60	64.3	5.00	1.16	10.3	S <sub>X</sub> 36.6(0.05) S <sub>C</sub> * 70.1(0.29) S <sub>A</sub> 75.8(0.79) I	I 69.0(-0.75) S <sub>A</sub> 66.5(-0.28) S <sub>C</sub> * 31.5(-0.07) S <sub>X</sub>	
					S <sub>X</sub> 35.4(0.08) S <sub>C</sub> * 70.1(0.29) S <sub>A</sub> 75.8(0.78) I		
50/50	83.8	5.18	1.15	10.6	S <sub>X</sub> 43.4(0.10) S <sub>C</sub> * 72.4(0.26) S <sub>A</sub> 77.3(0.85) I	I 70.4(-0.84) S <sub>A</sub> 68.6(-0.31) S <sub>C</sub> * 38.9(-0.12) S <sub>X</sub>	
					S <sub>X</sub> 43.0(0.13) S <sub>C</sub> * 72.4(0.27) S <sub>A</sub> 77.2(0.90) I		
60/40	81.7	5.20	1.15	10.7	S <sub>X</sub> 42.1(0.09) S <sub>C</sub> * 72.0(0.25) S <sub>A</sub> 76.8(0.84) I	I 70.2(-0.75) S <sub>A</sub> 68.4(-0.28) S <sub>C</sub> * 37.7(-0.10) S <sub>X</sub>	
					S <sub>X</sub> 41.5(0.10) S <sub>C</sub> * 71.8(0.26) S <sub>A</sub> 76.5(0.82) I		
80/20	78.8	4.96	1.17	10.2	S <sub>X</sub> 39.3(0.06) S <sub>C</sub> * 70.6(0.30) S <sub>A</sub> 76.0(0.81) I	I 68.3(-1.12) S <sub>C</sub> * 33.0(-0.07) S <sub>X</sub>	
					S <sub>X</sub> 38.2(0.10) S <sub>C</sub> * 70.5(0.32) S <sub>A</sub> 75.6(0.86) I		
100/0	91.0	4.98	1.12	10.2	K 37.3(0.25) S <sub>X</sub> 40.0(† <sup>c</sup> ) S <sub>C</sub> * 72.6(0.29) S <sub>A</sub> 77.0(0.78) I	I 70.2(-0.97) S <sub>C</sub> * 34.9(-0.06) S <sub>X</sub>	
					S <sub>X</sub> 39.3(0.08) S <sub>C</sub> * 72.4(0.29) S <sub>A</sub> 76.5(0.79) I		

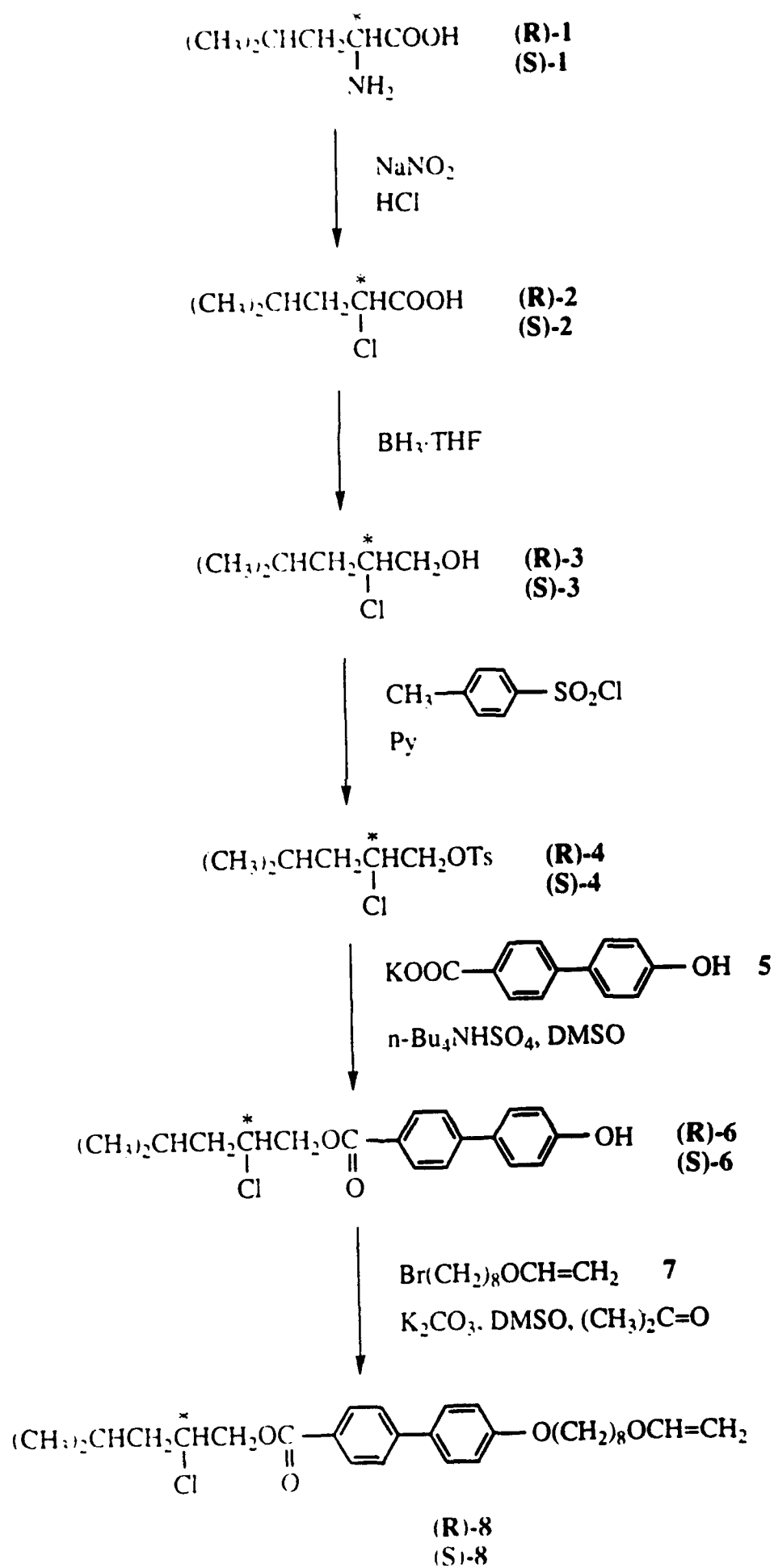
<sup>a</sup> Polymerization temperature: 0°C; polymerization solvent: methylene chloride; [M]<sub>0</sub>=[(R)-8]+[(S)-8]=0.224;

[M]<sub>0</sub>/[I]<sub>0</sub>=16; [Me<sub>2</sub>S]<sub>0</sub>/[I]<sub>0</sub>=10; polymerization time: 1 hour.

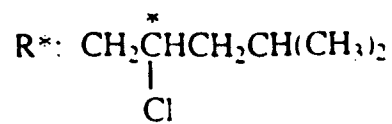
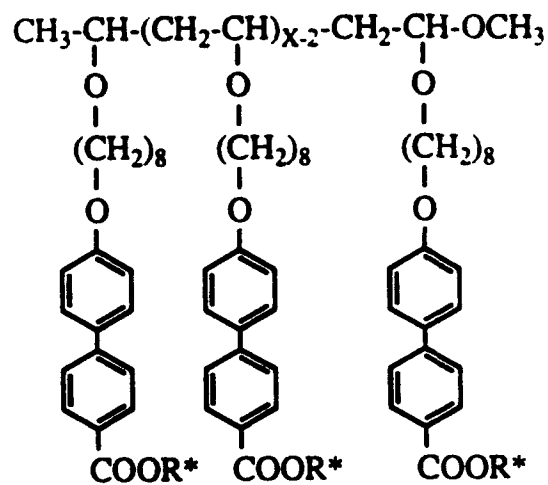
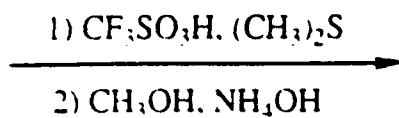
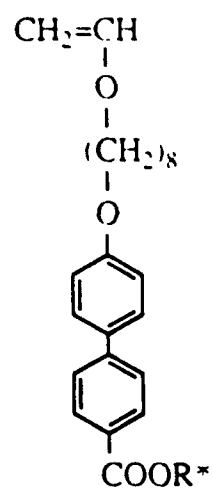
<sup>b</sup> Heating and cooling rates are 10°C/min.

Data on the first line are from first heating and cooling scans. Data on the second line are from second heating scan.

<sup>c</sup> Overlapped peak.



Scheme 1



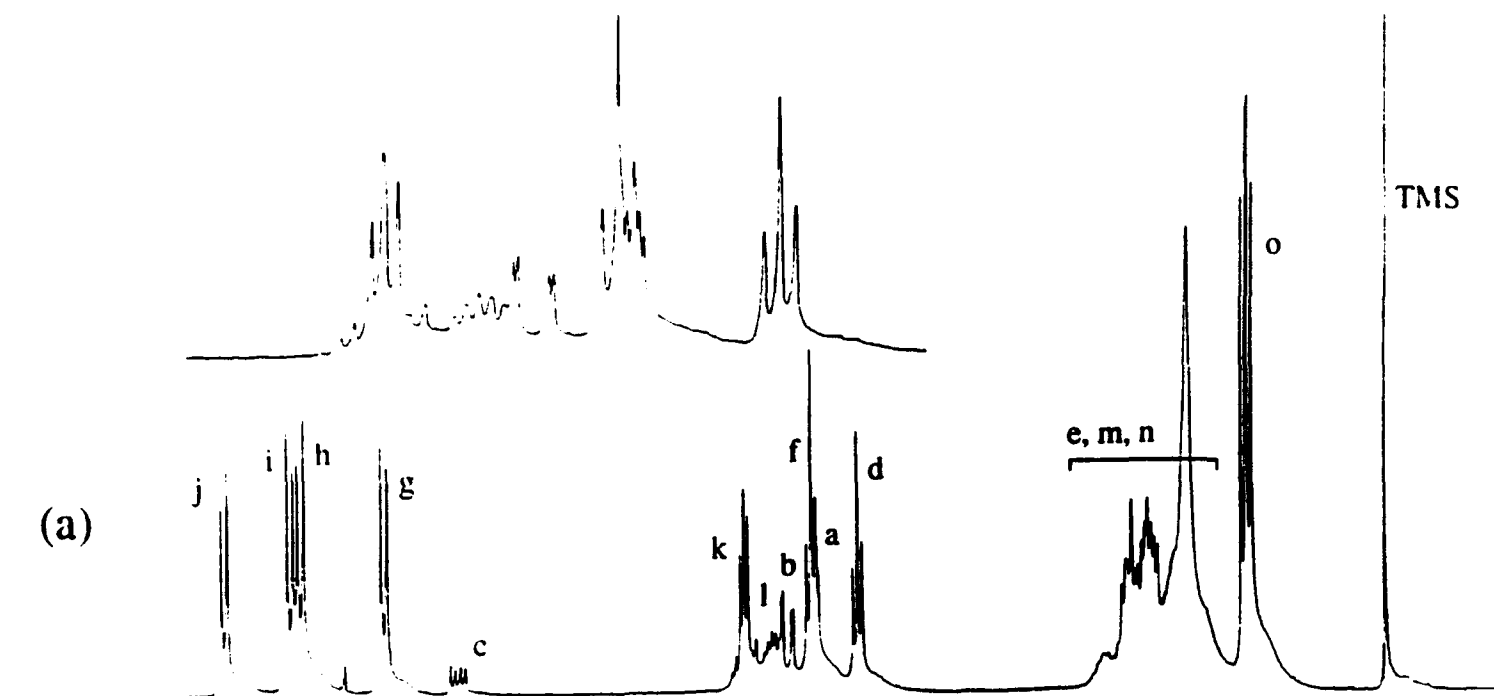
(R)-8

(S)-8

Scheme 2



Scheme 3





10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100 101 102 103 104 105 106 107 108 109 110 111 112 113 114 115 116 117 118 119 120 121 122 123 124 125 126 127 128 129 130 131 132 133 134 135 136 137 138 139 140 141 142 143 144 145 146 147 148 149 150 151 152 153 154 155 156 157 158 159 160 161 162 163 164 165 166 167 168 169 170 171 172 173 174 175 176 177 178 179 180 181 182 183 184 185 186 187 188 189 190 191 192 193 194 195 196 197 198 199 200 201 202 203 204 205 206 207 208 209 210 211 212 213 214 215 216 217 218 219 220 221 222 223 224 225 226 227 228 229 230 231 232 233 234 235 236 237 238 239 240 241 242 243 244 245 246 247 248 249 250 251 252 253 254 255 256 257 258 259 260 261 262 263 264 265 266 267 268 269 270 271 272 273 274 275 276 277 278 279 280 281 282 283 284 285 286 287 288 289 290 291 292 293 294 295 296 297 298 299 300 301 302 303 304 305 306 307 308 309 310 311 312 313 314 315 316 317 318 319 320 321 322 323 324 325 326 327 328 329 330 331 332 333 334 335 336 337 338 339 340 341 342 343 344 345 346 347 348 349 350 351 352 353 354 355 356 357 358 359 360 361 362 363 364 365 366 367 368 369 370 371 372 373 374 375 376 377 378 379 380 381 382 383 384 385 386 387 388 389 390 391 392 393 394 395 396 397 398 399 400 401 402 403 404 405 406 407 408 409 410 411 412 413 414 415 416 417 418 419 420 421 422 423 424 425 426 427 428 429 430 431 432 433 434 435 436 437 438 439 440 441 442 443 444 445 446 447 448 449 450 451 452 453 454 455 456 457 458 459 460 461 462 463 464 465 466 467 468 469 470 471 472 473 474 475 476 477 478 479 480 481 482 483 484 485 486 487 488 489 490 491 492 493 494 495 496 497 498 499 500 501 502 503 504 505 506 507 508 509 510 511 512 513 514 515 516 517 518 519 520 521 522 523 524 525 526 527 528 529 530 531 532 533 534 535 536 537 538 539 540 541 542 543 544 545 546 547 548 549 550 551 552 553 554 555 556 557 558 559 560 561 562 563 564 565 566 567 568 569 570 571 572 573 574 575 576 577 578 579 580 581 582 583 584 585 586 587 588 589 590 591 592 593 594 595 596 597 598 599 600 601 602 603 604 605 606 607 608 609 610 611 612 613 614 615 616 617 618 619 620 621 622 623 624 625 626 627 628 629 630 631 632 633 634 635 636 637 638 639 640 641 642 643 644 645 646 647 648 649 650 651 652 653 654 655 656 657 658 659 660 661 662 663 664 665 666 667 668 669 670 671 672 673 674 675 676 677 678 679 680 681 682 683 684 685 686 687 688 689 690 691 692 693 694 695 696 697 698 699 700 701 702 703 704 705 706 707 708 709 710 711 712 713 714 715 716 717 718 719 720 721 722 723 724 725 726 727 728 729 730 731 732 733 734 735 736 737 738 739 740 741 742 743 744 745 746 747 748 749 750 751 752 753 754 755 756 757 758 759 760 761 762 763 764 765 766 767 768 769 770 771 772 773 774 775 776 777 778 779 780 781 782 783 784 785 786 787 788 789 790 791 792 793 794 795 796 797 798 799 800 801 802 803 804 805 806 807 808 809 810 811 812 813 814 815 816 817 818 819 820 821 822 823 824 825 826 827 828 829 830 831 832 833 834 835 836 837 838 839 840 841 842 843 844 845 846 847 848 849 850 851 852 853 854 855 856 857 858 859 860 861 862 863 864 865 866 867 868 869 870 871 872 873 874 875 876 877 878 879 880 881 882 883 884 885 886 887 888 889 890 891 892 893 894 895 896 897 898 899 900 901 902 903 904 905 906 907 908 909 910 911 912 913 914 915 916 917 918 919 920 921 922 923 924 925 926 927 928 929 930 931 932 933 934 935 936 937 938 939 940 941 942 943 944 945 946 947 948 949 950 951 952 953 954 955 956 957 958 959 960 961 962 963 964 965 966 967 968 969 970 971 972 973 974 975 976 977 978 979 980 981 982 983 984 985 986 987 988 989 990 991 992 993 994 995 996 997 998 999 1000 1001 1002 1003 1004 1005 1006 1007 1008 1009 1010 1011 1012 1013 1014 1015 1016 1017 1018 1019 1020 1021 1022 1023 1024 1025 1026 1027 1028 1029 1030 1031 1032 1033 1034 1035 1036 1037 1038 1039 1040 1041 1042 1043 1044

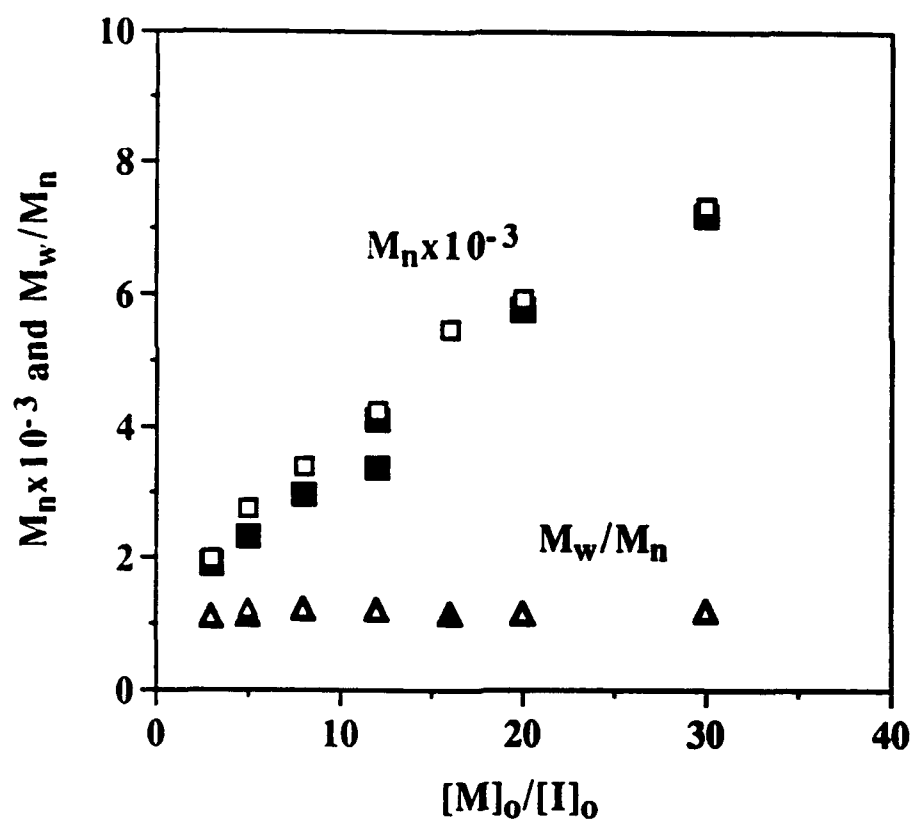
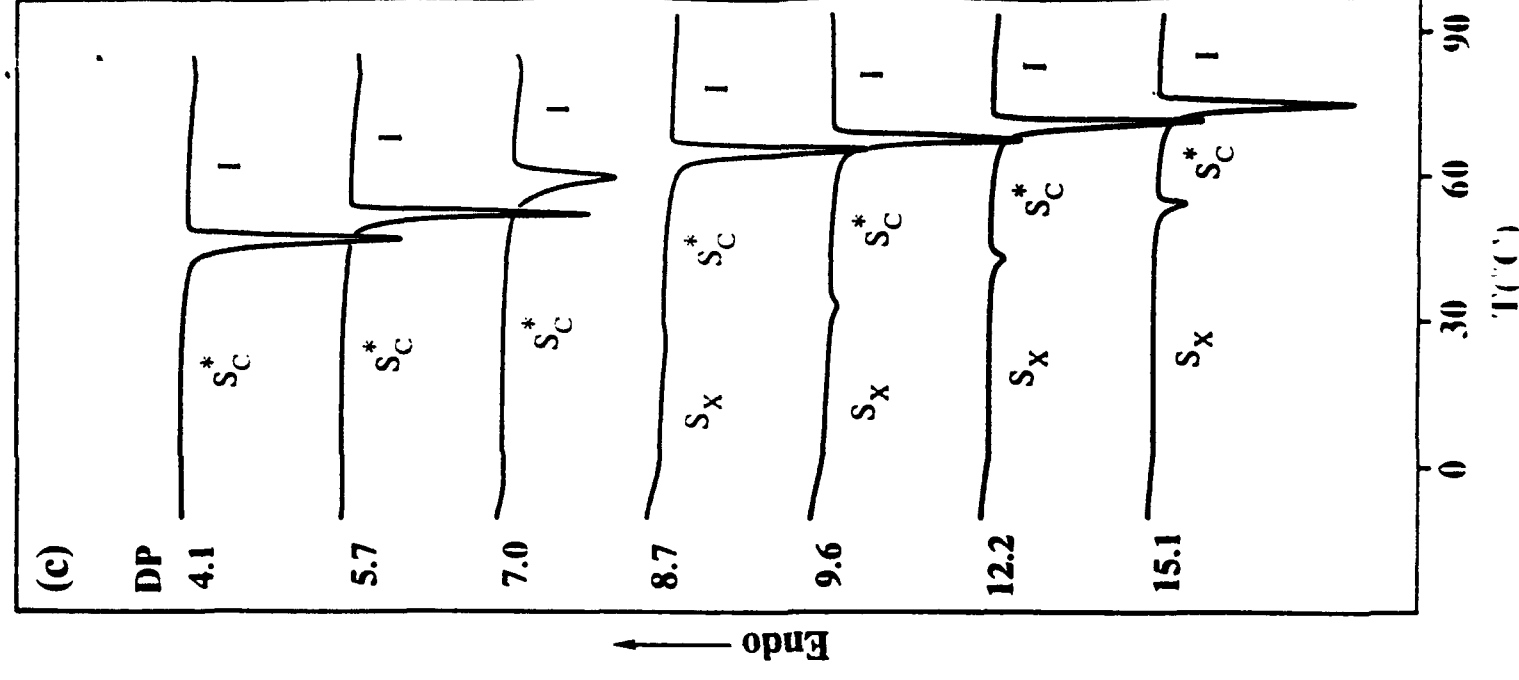
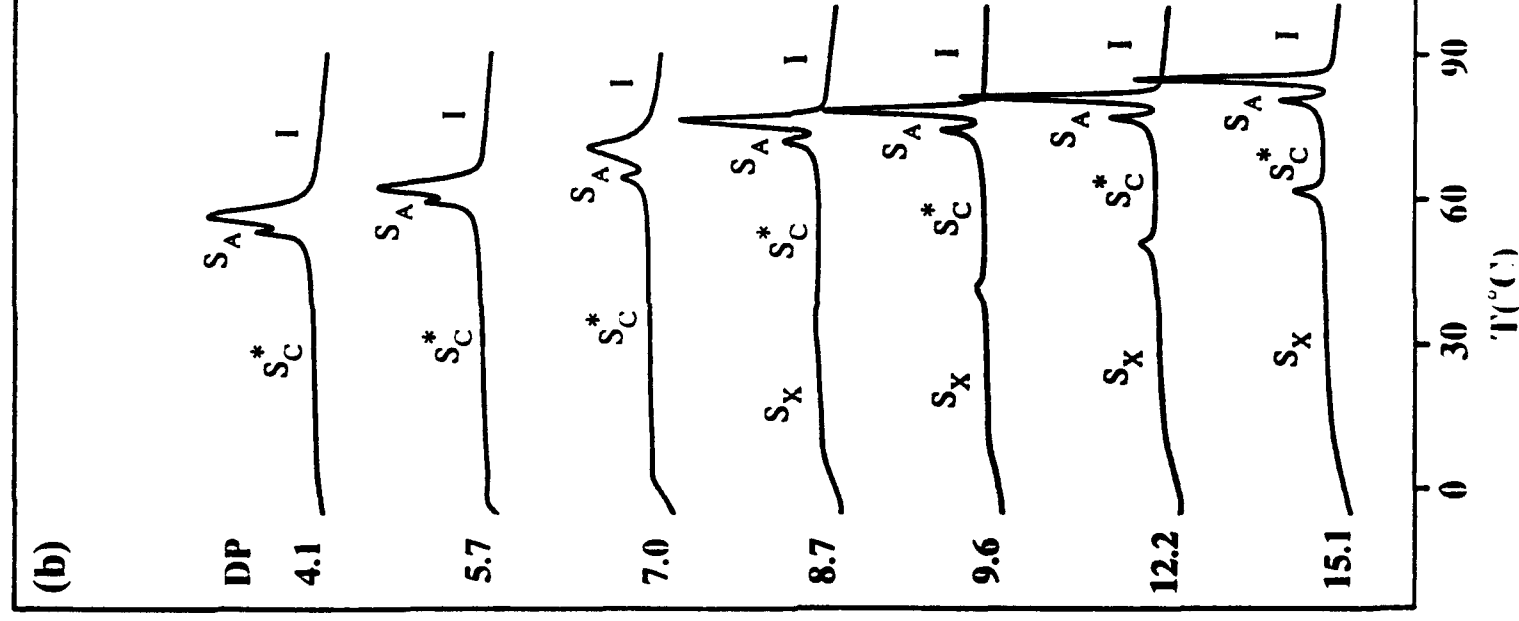
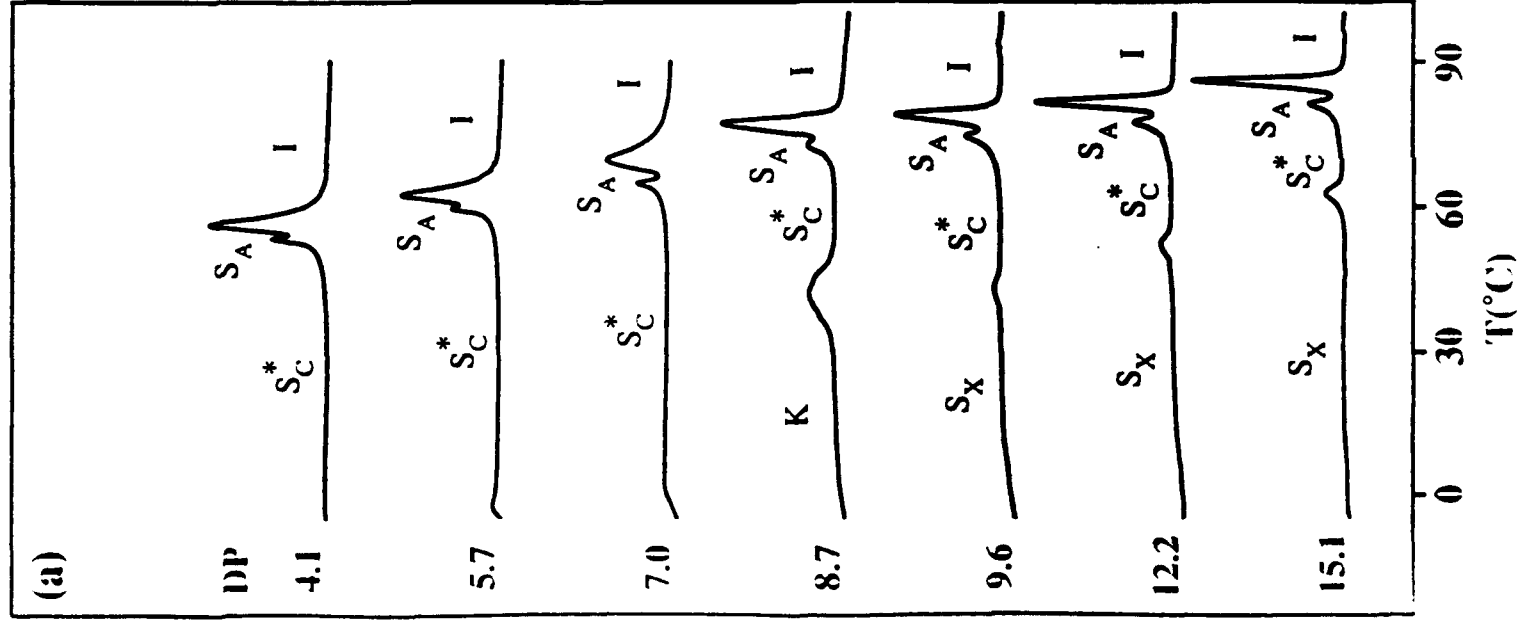
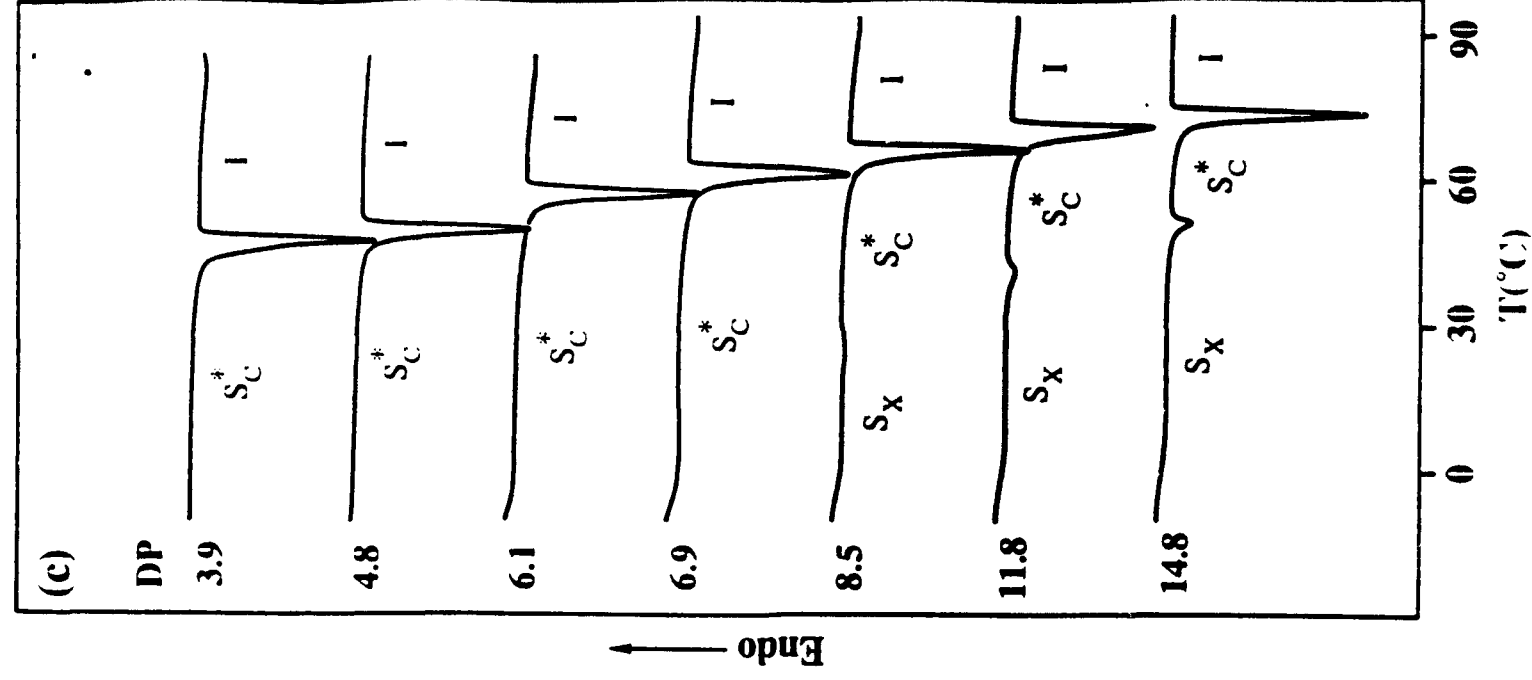
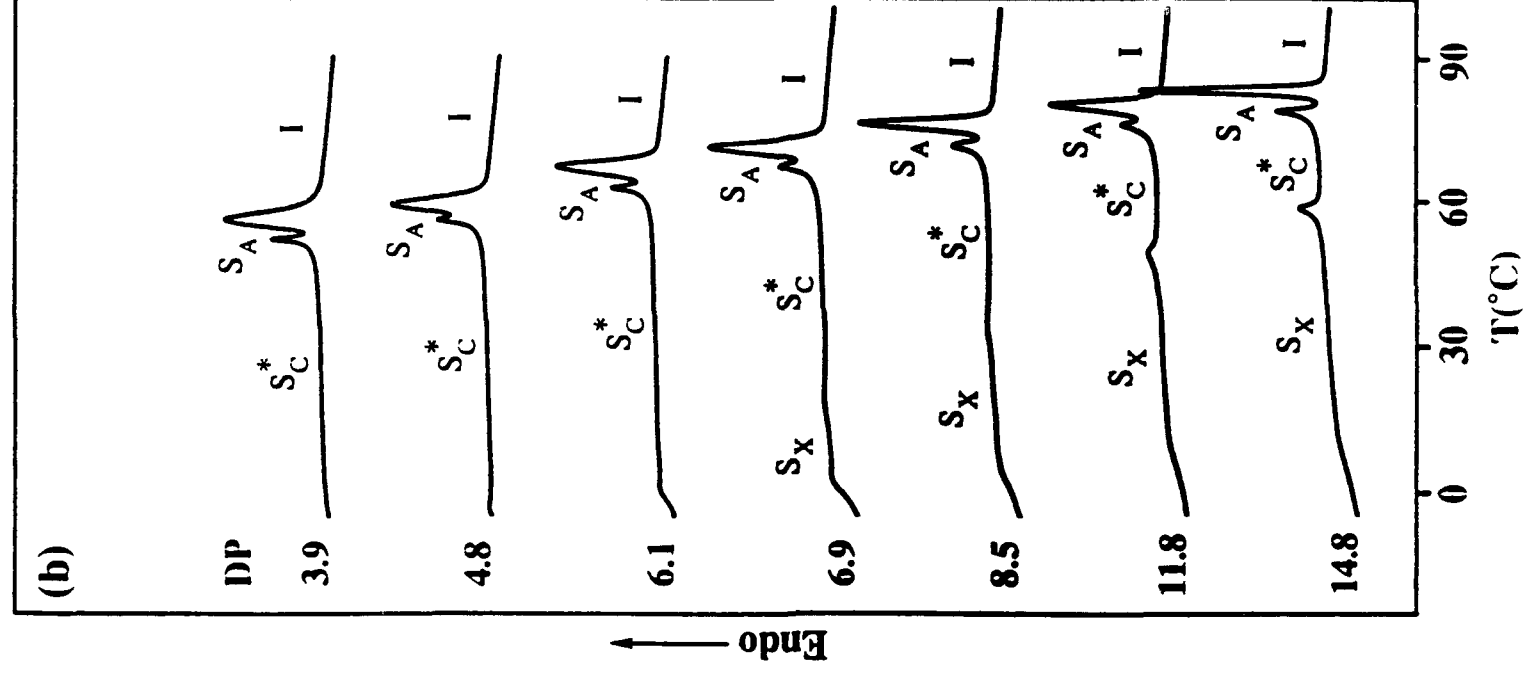
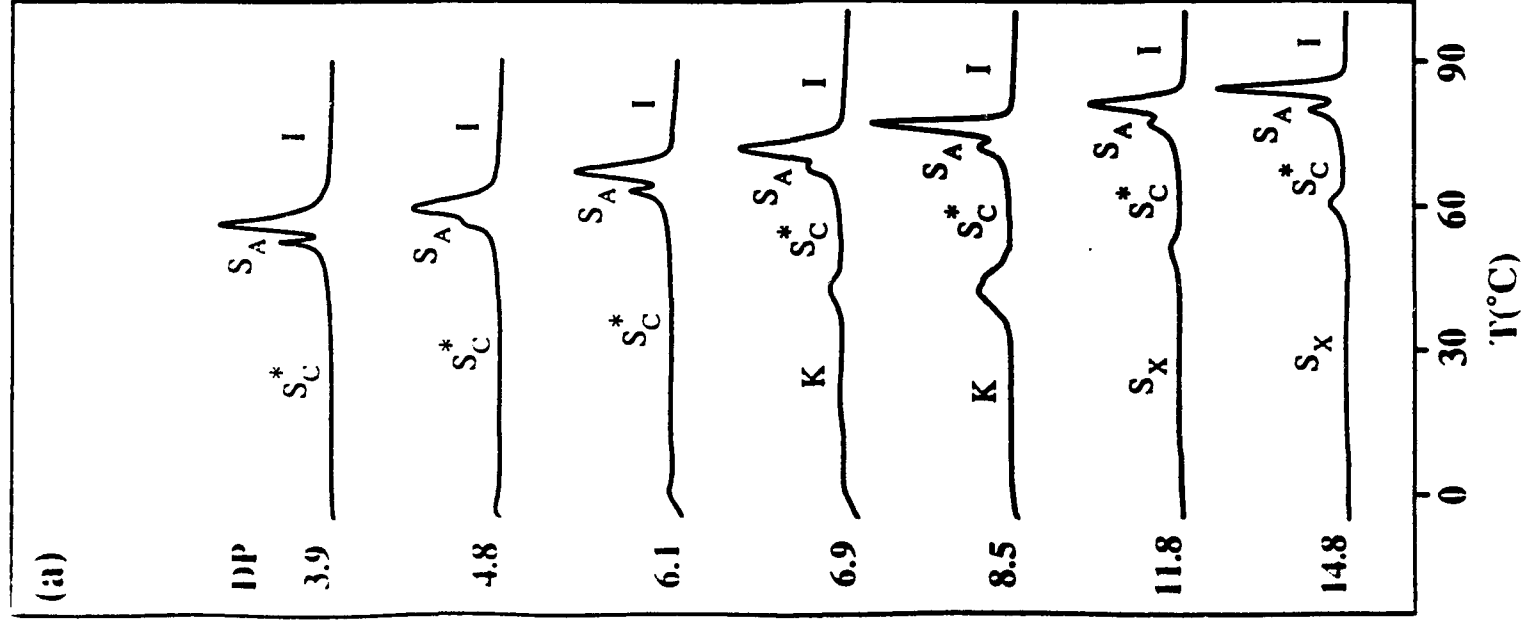


Figure 3







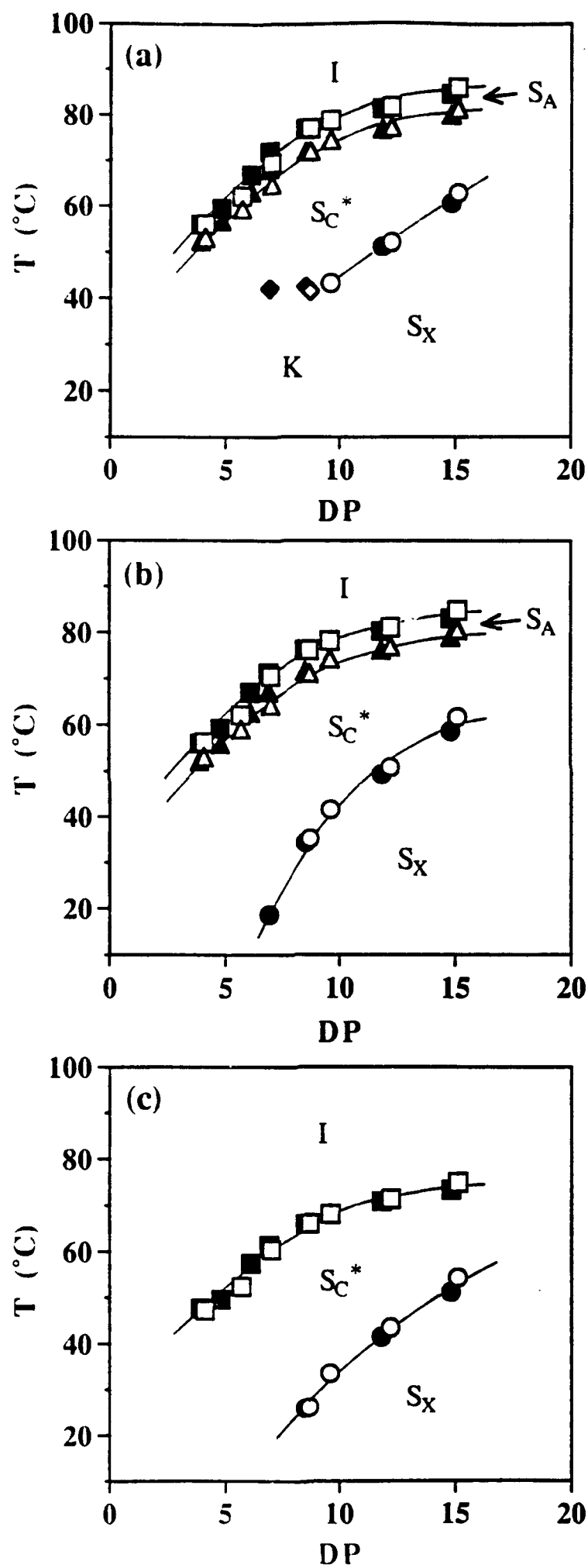
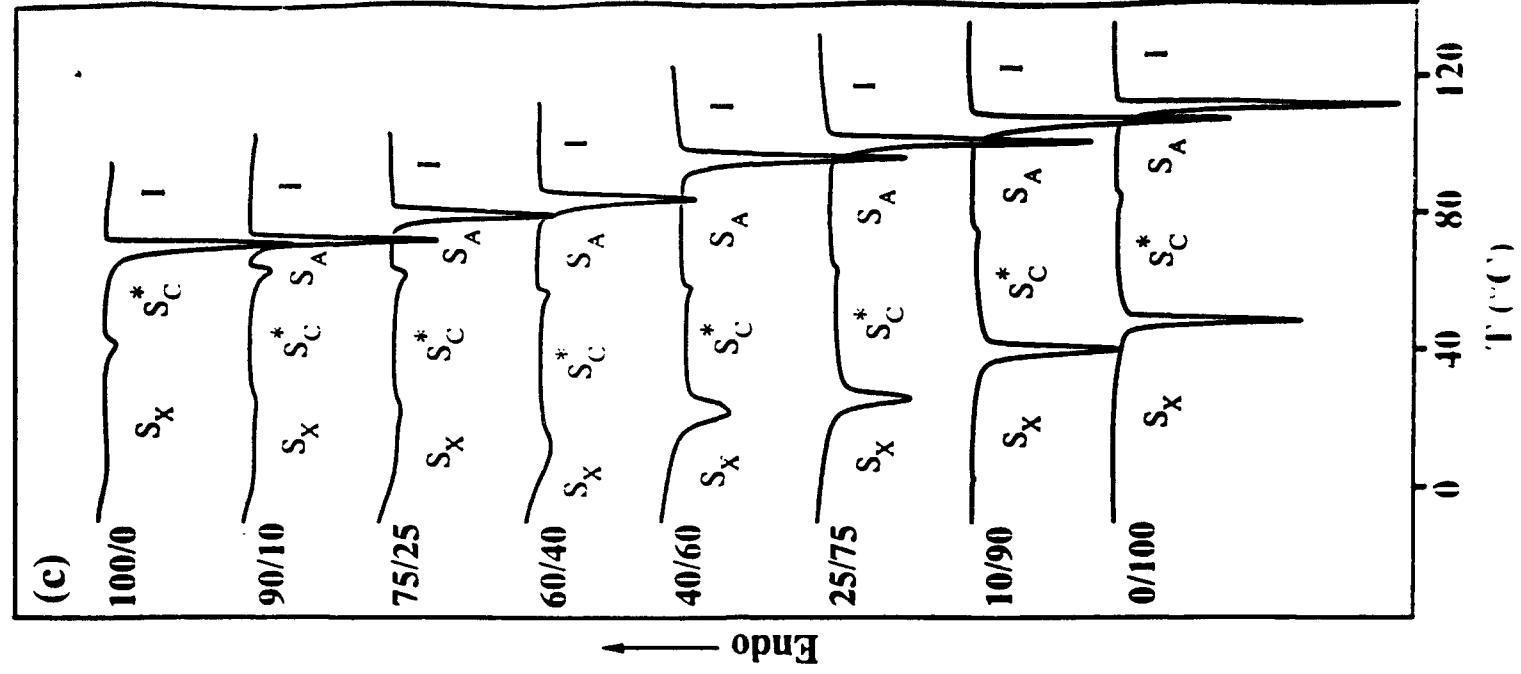
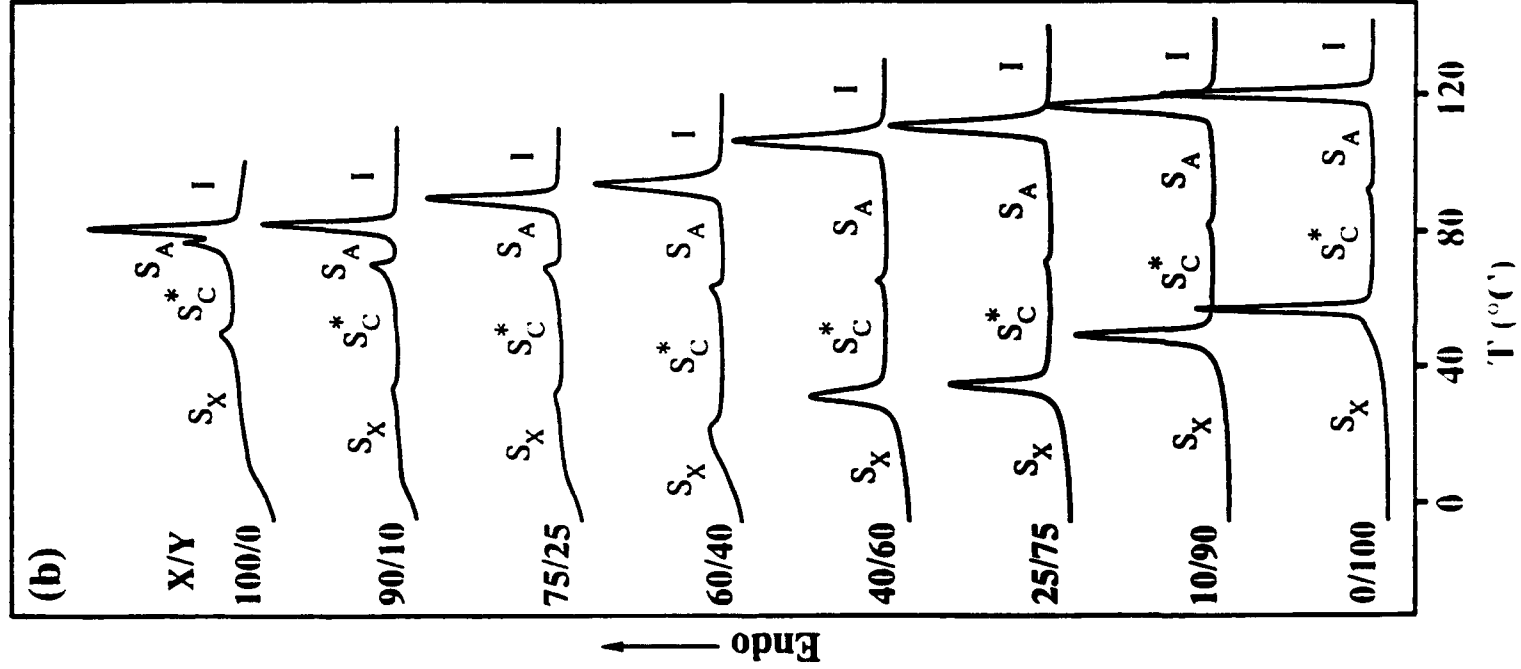
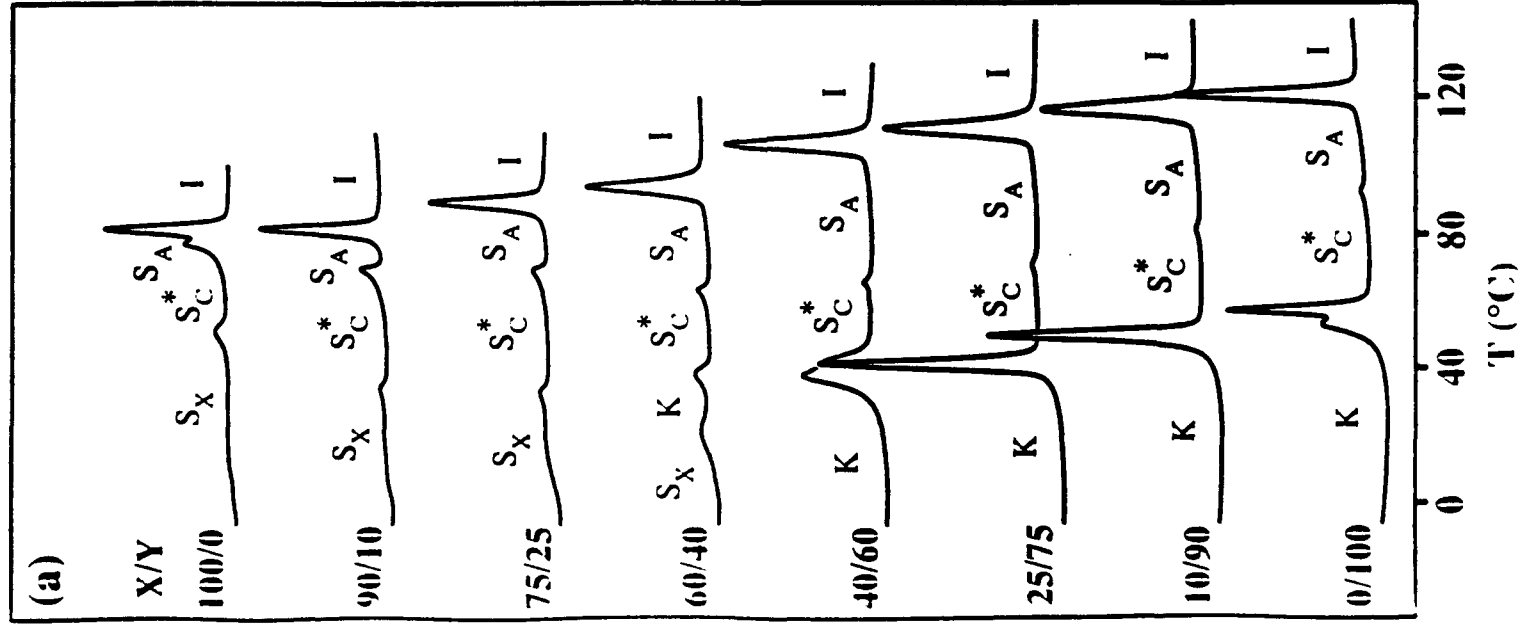


Figure 6



Figure 7



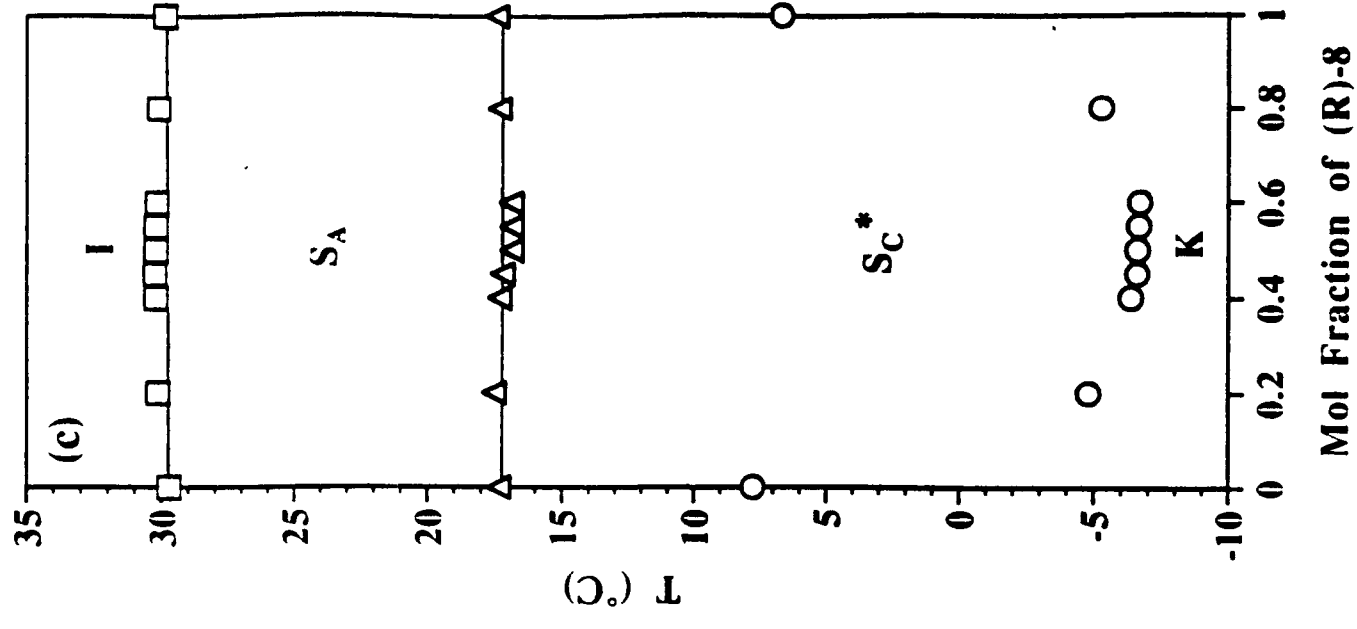
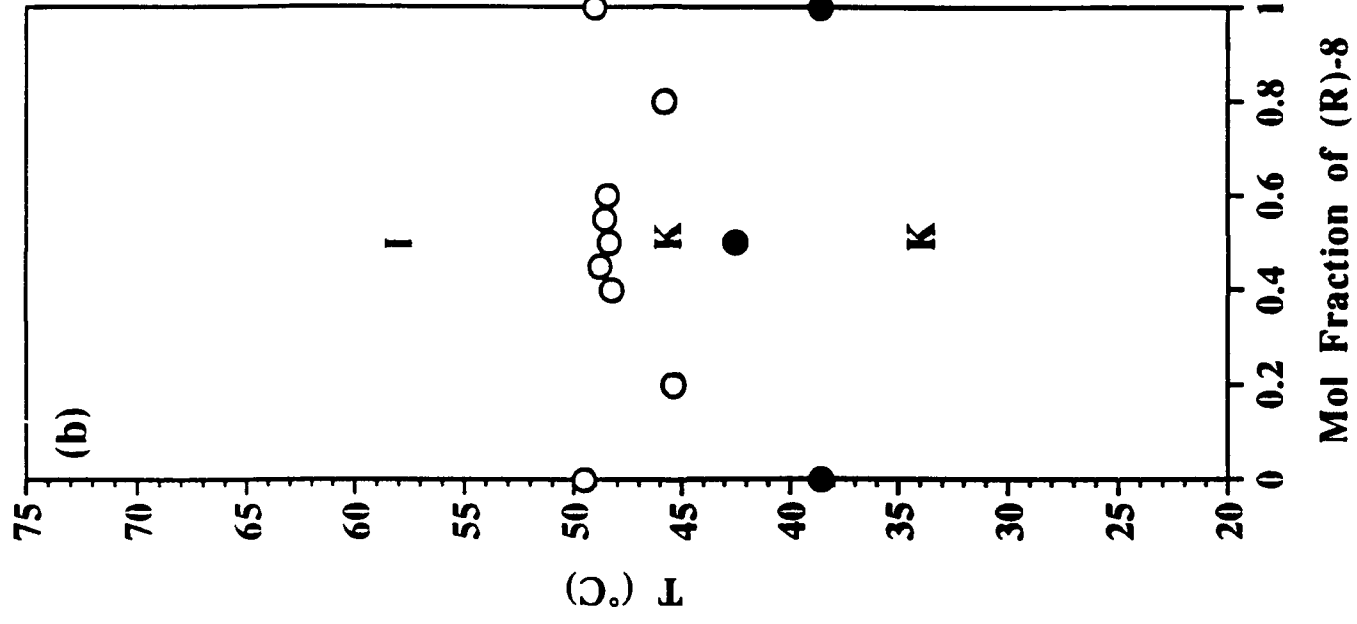
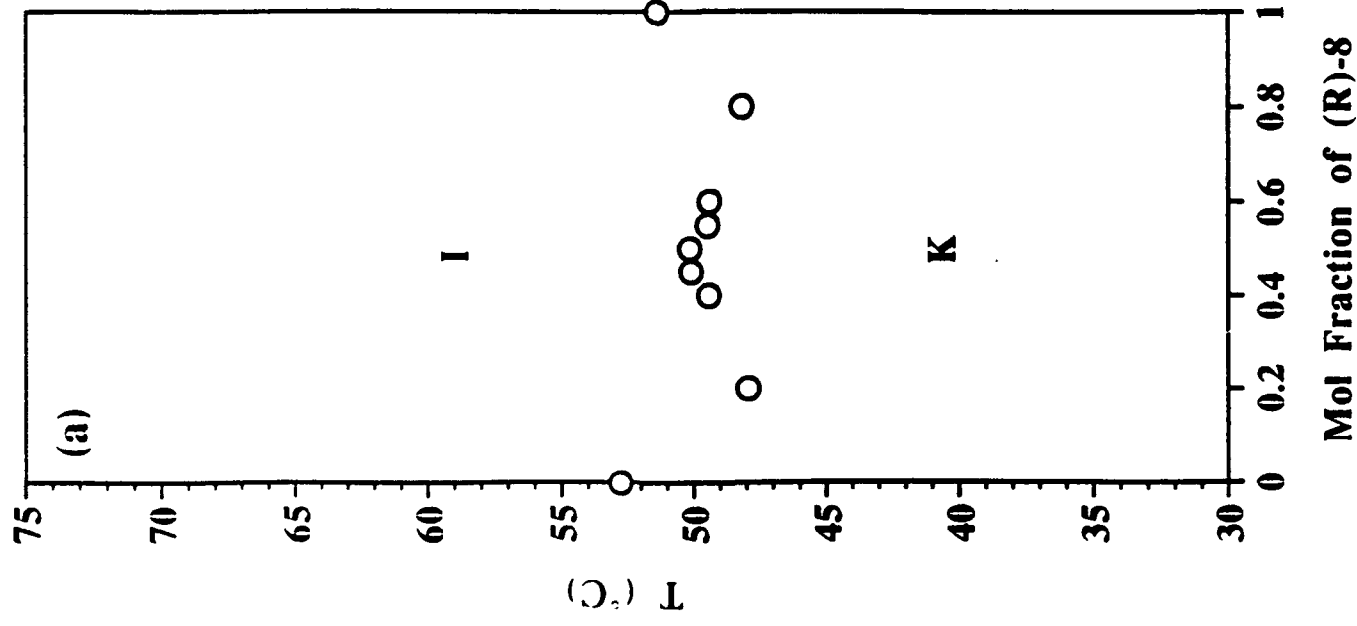


Figure 10

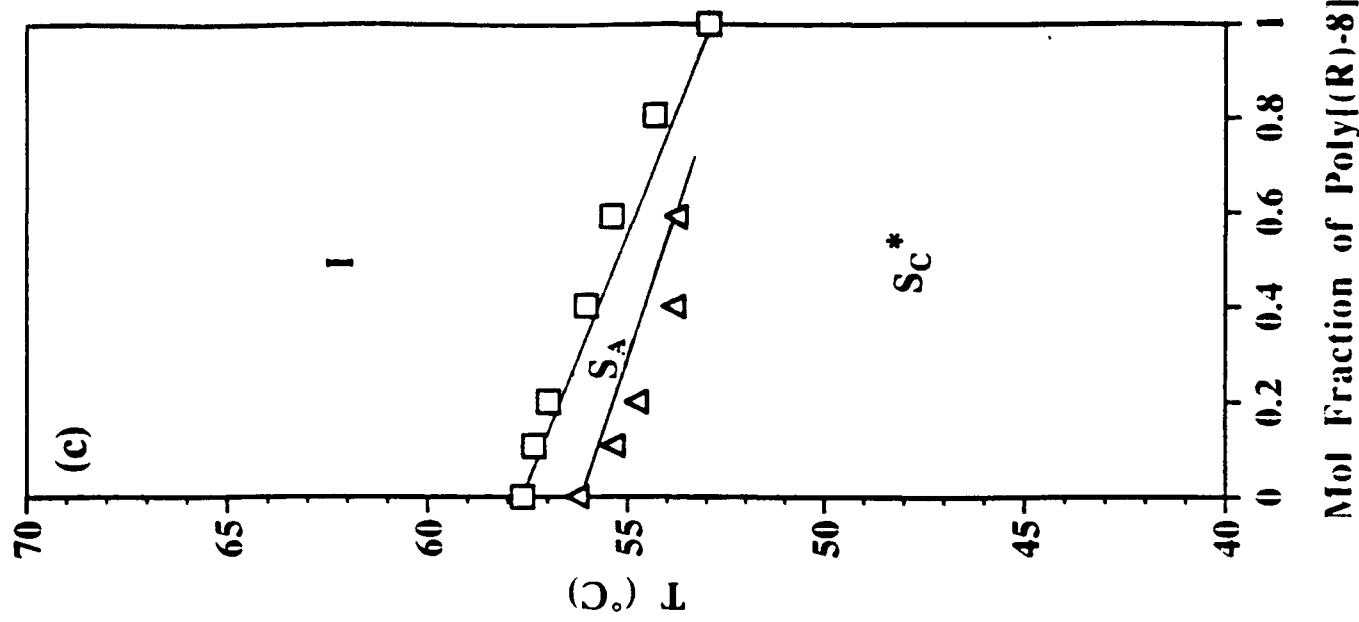
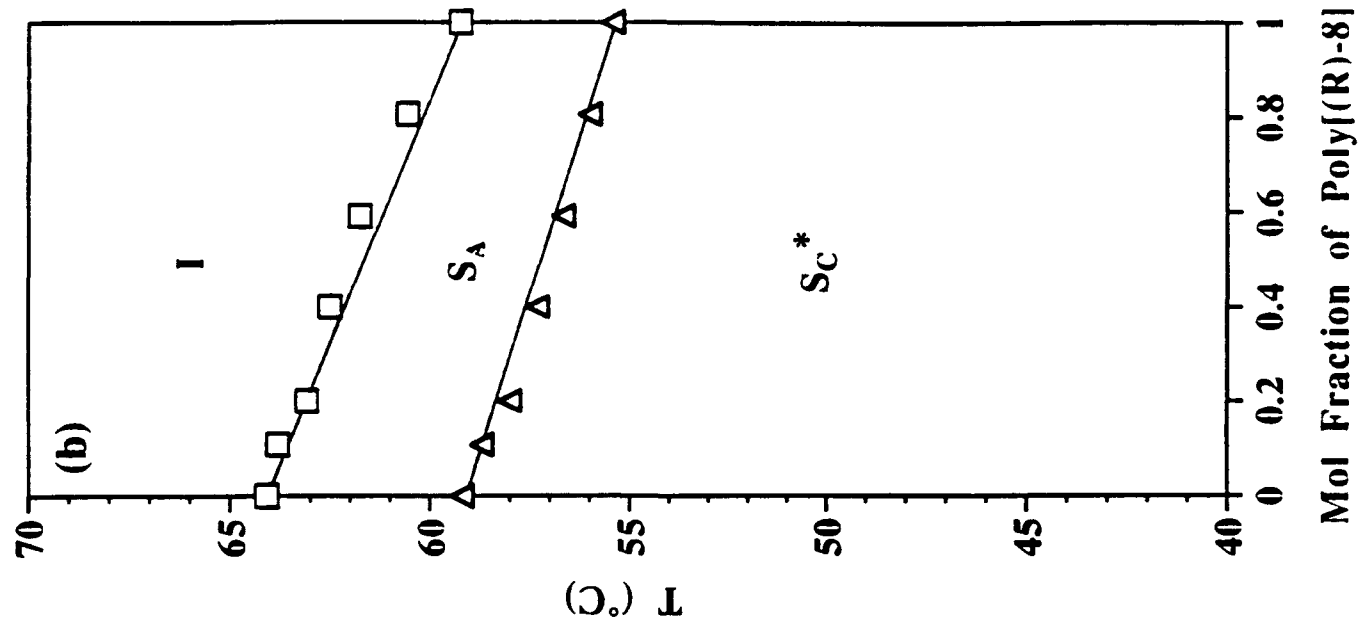
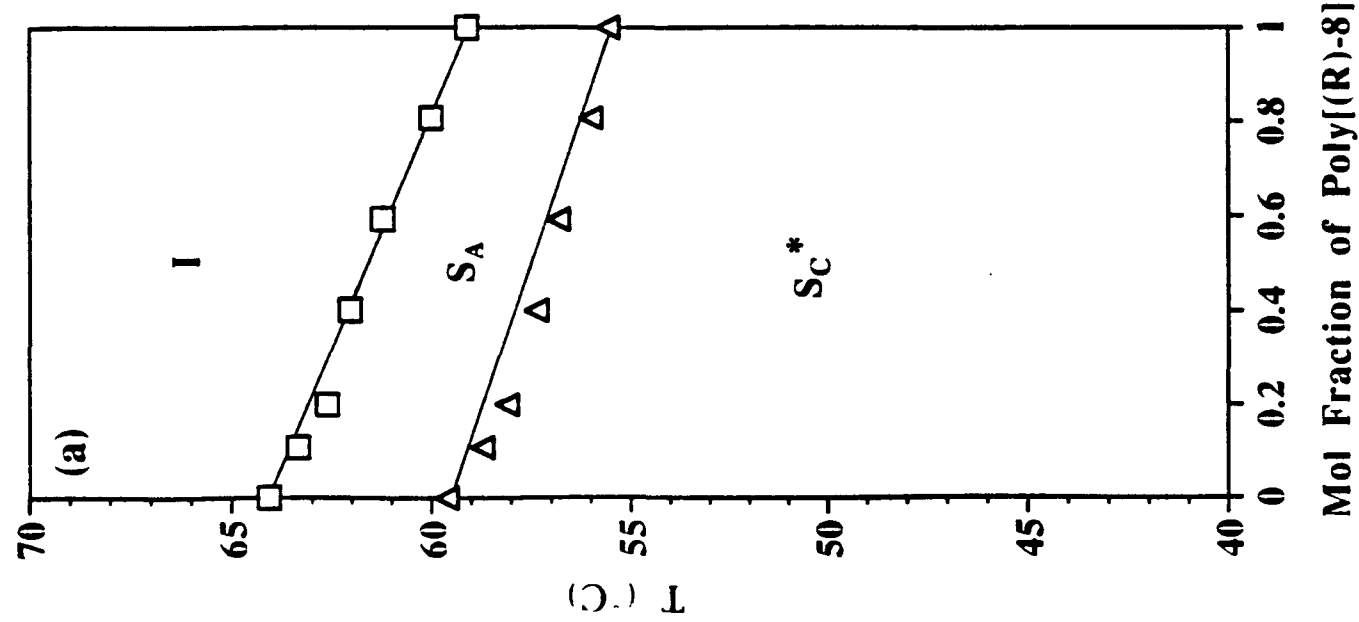


Figure 11

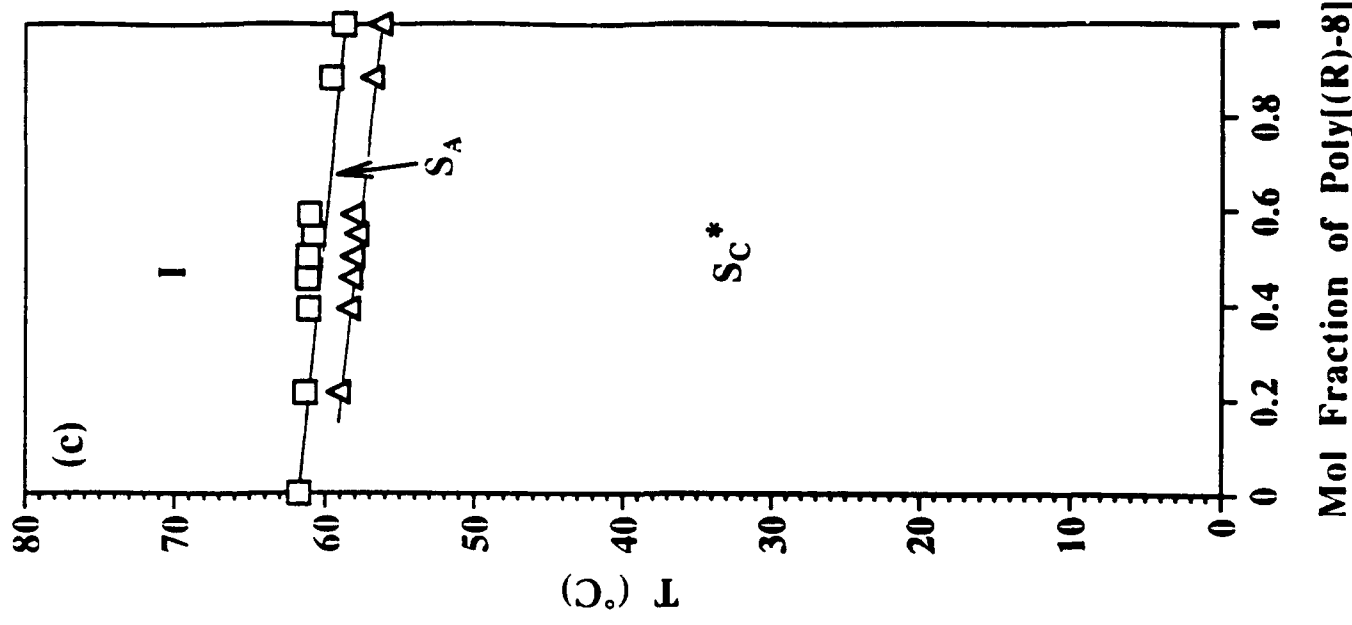
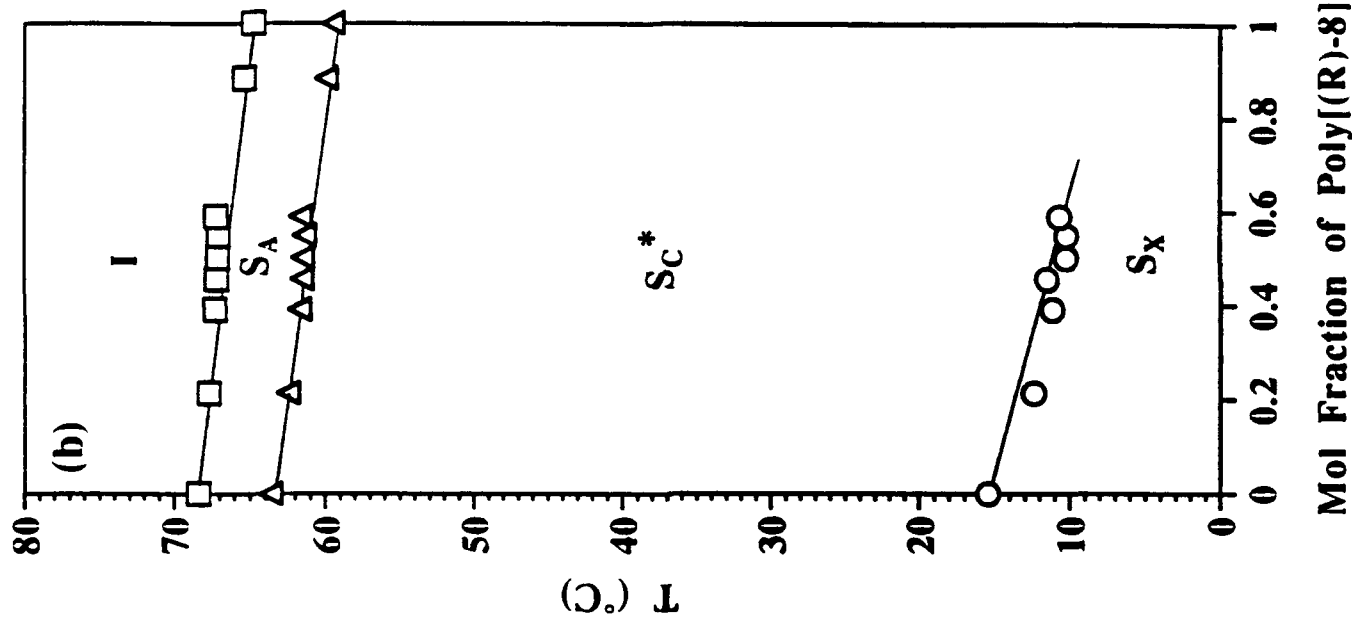
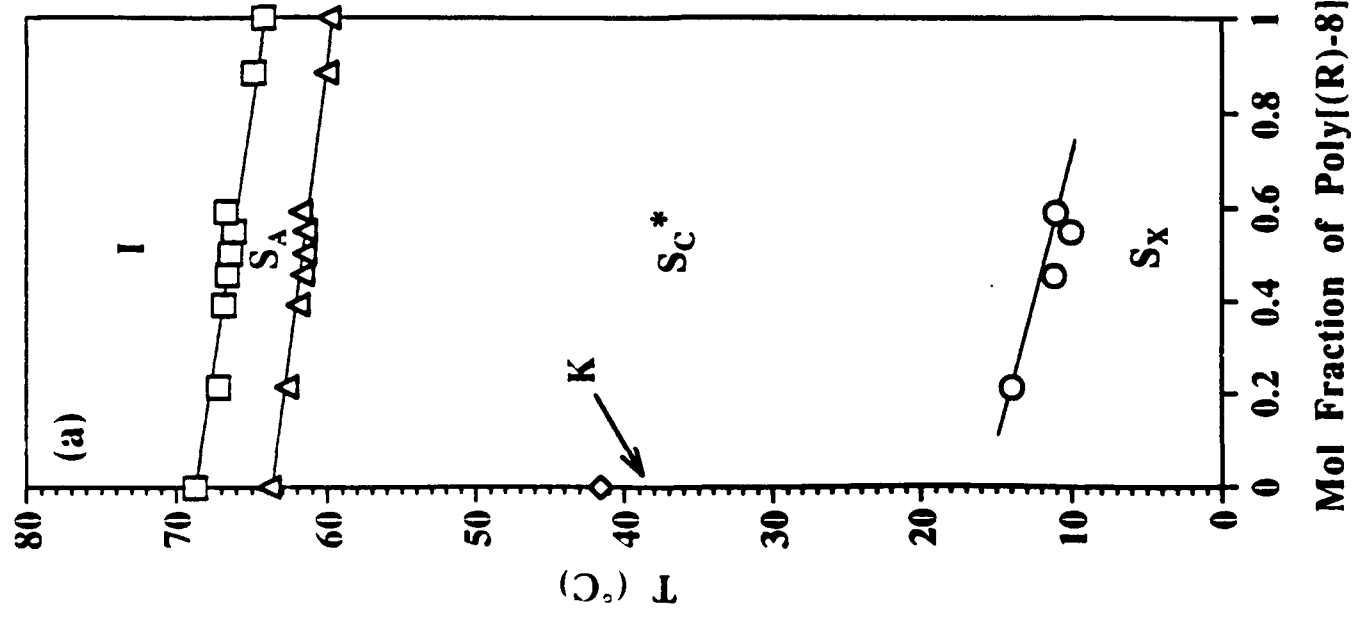


Figure 12

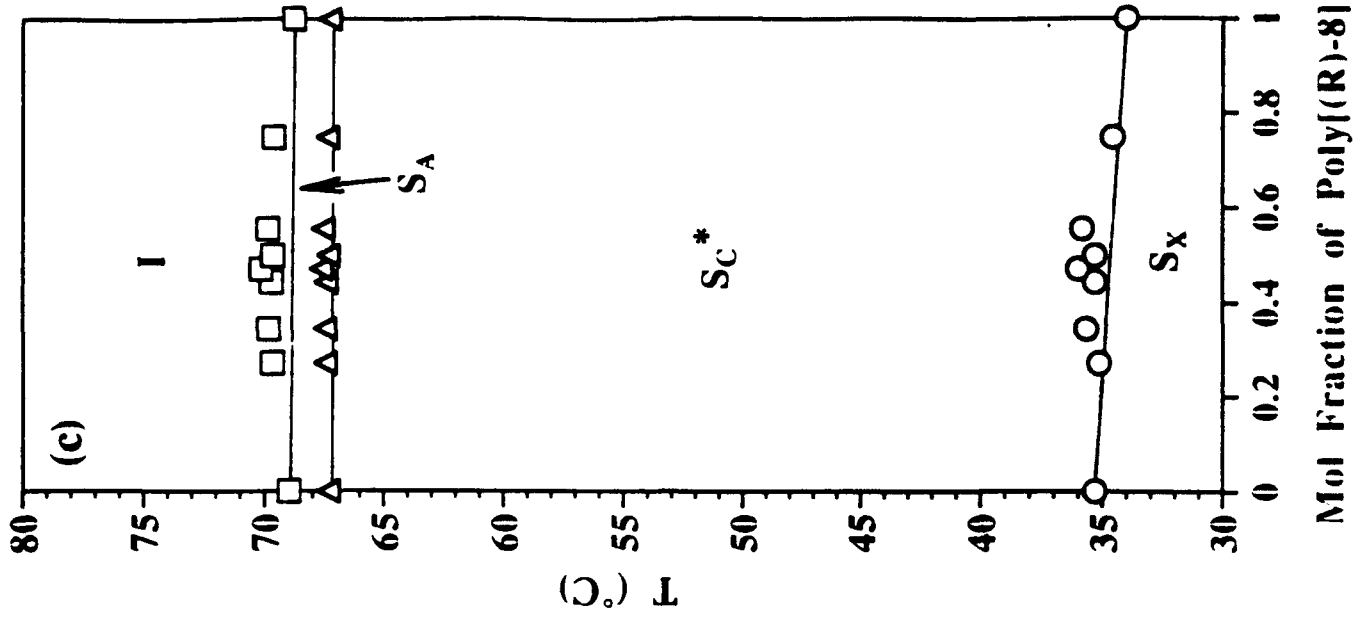
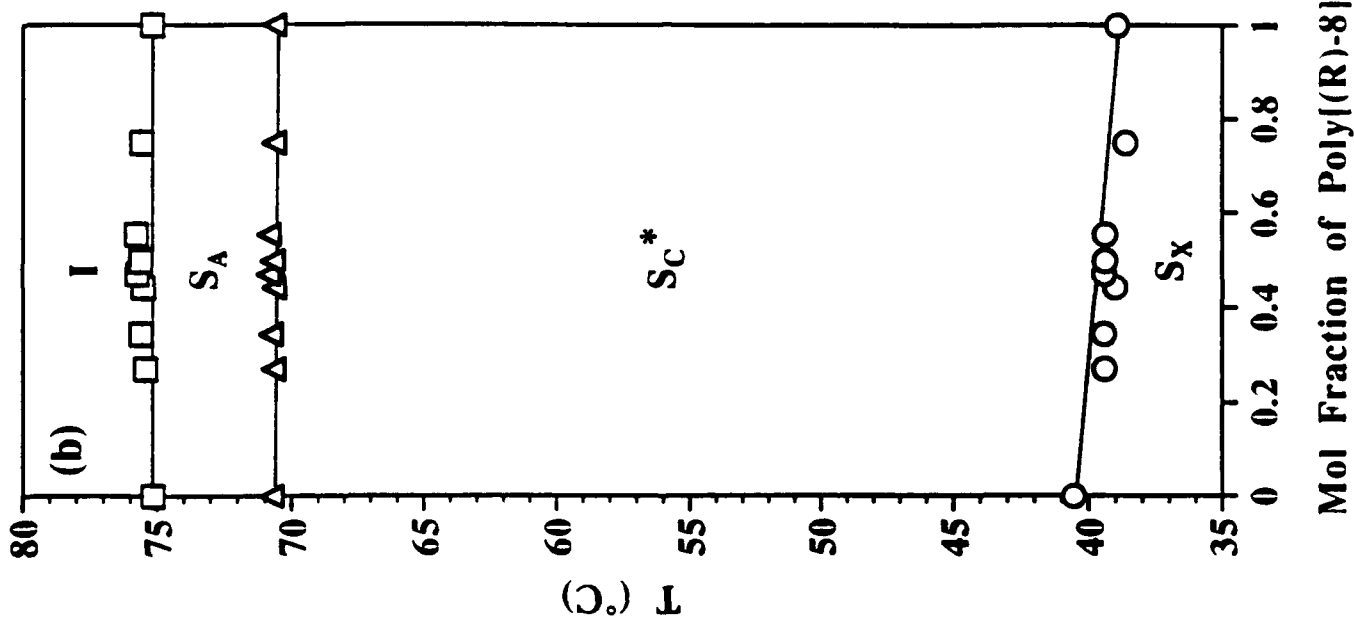
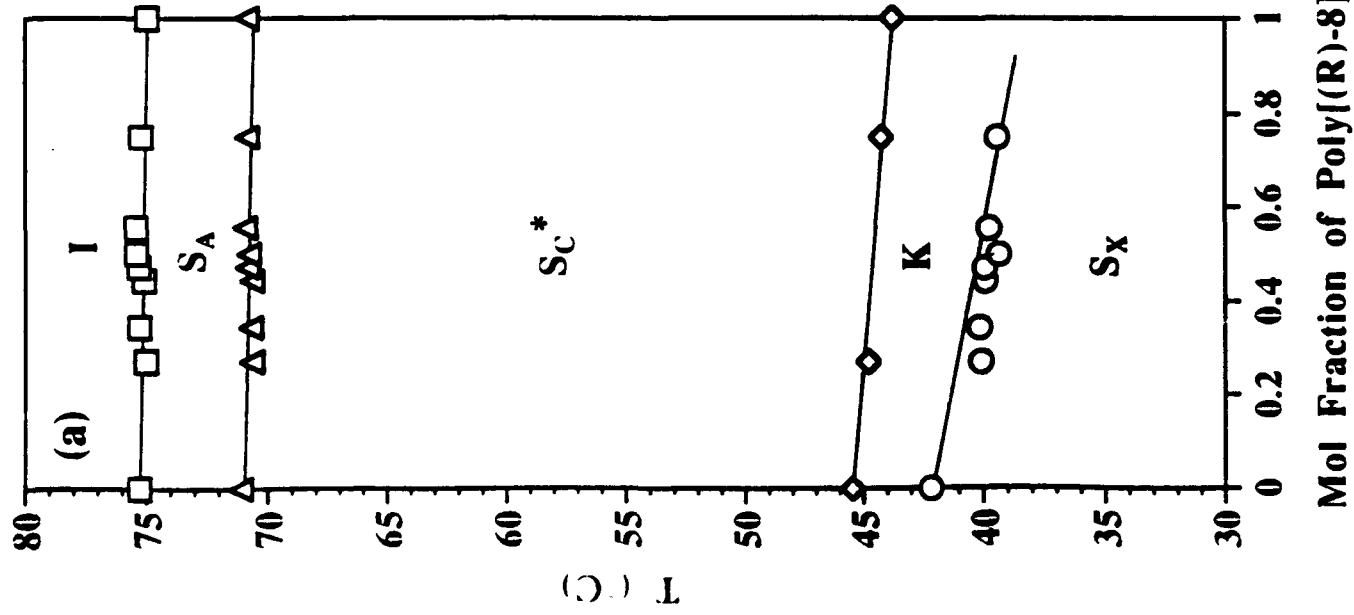


Figure 13



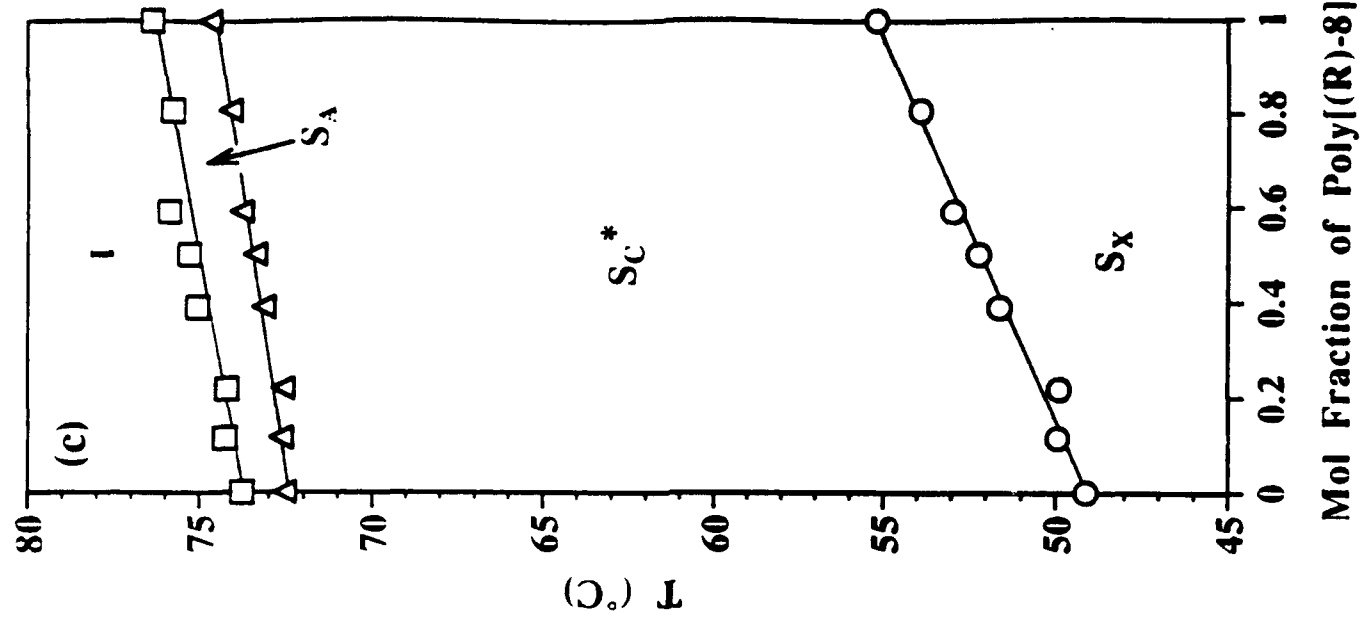
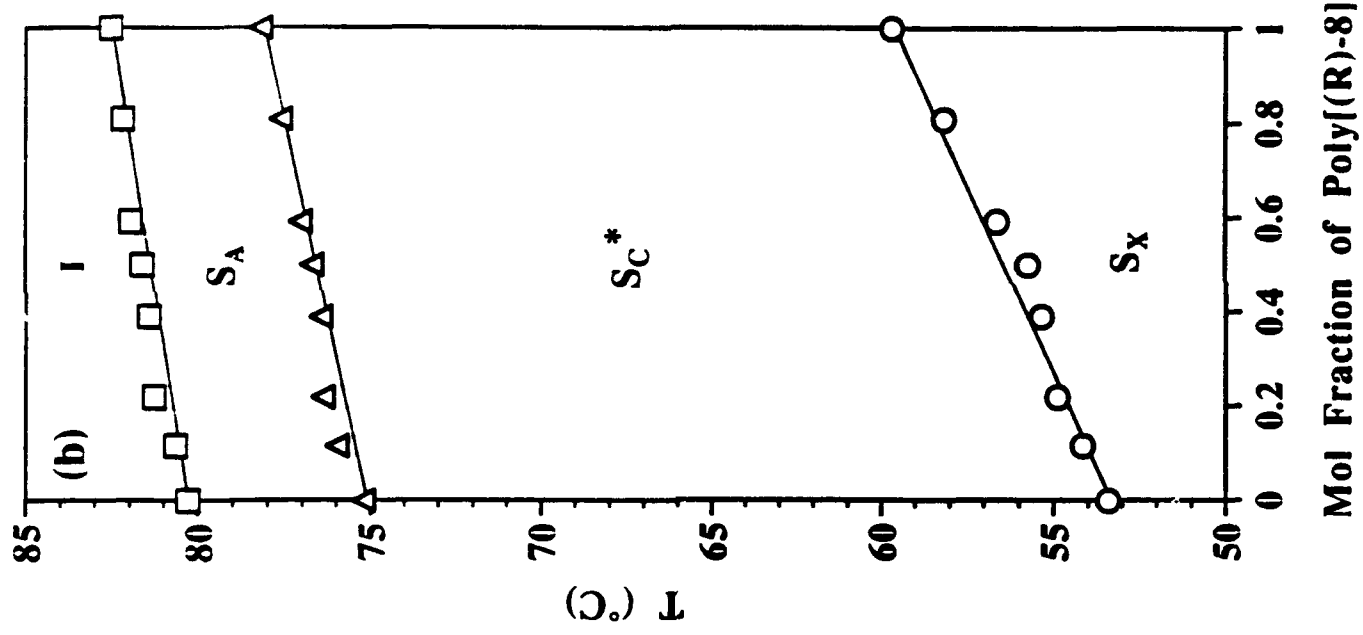
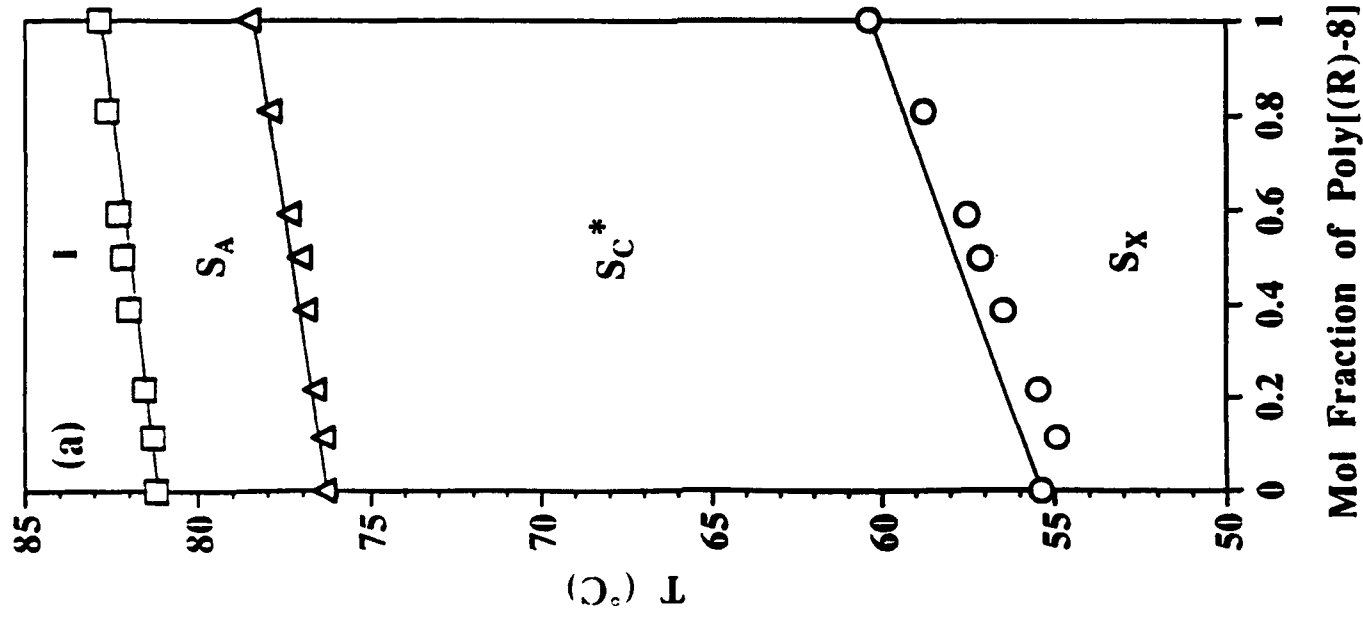
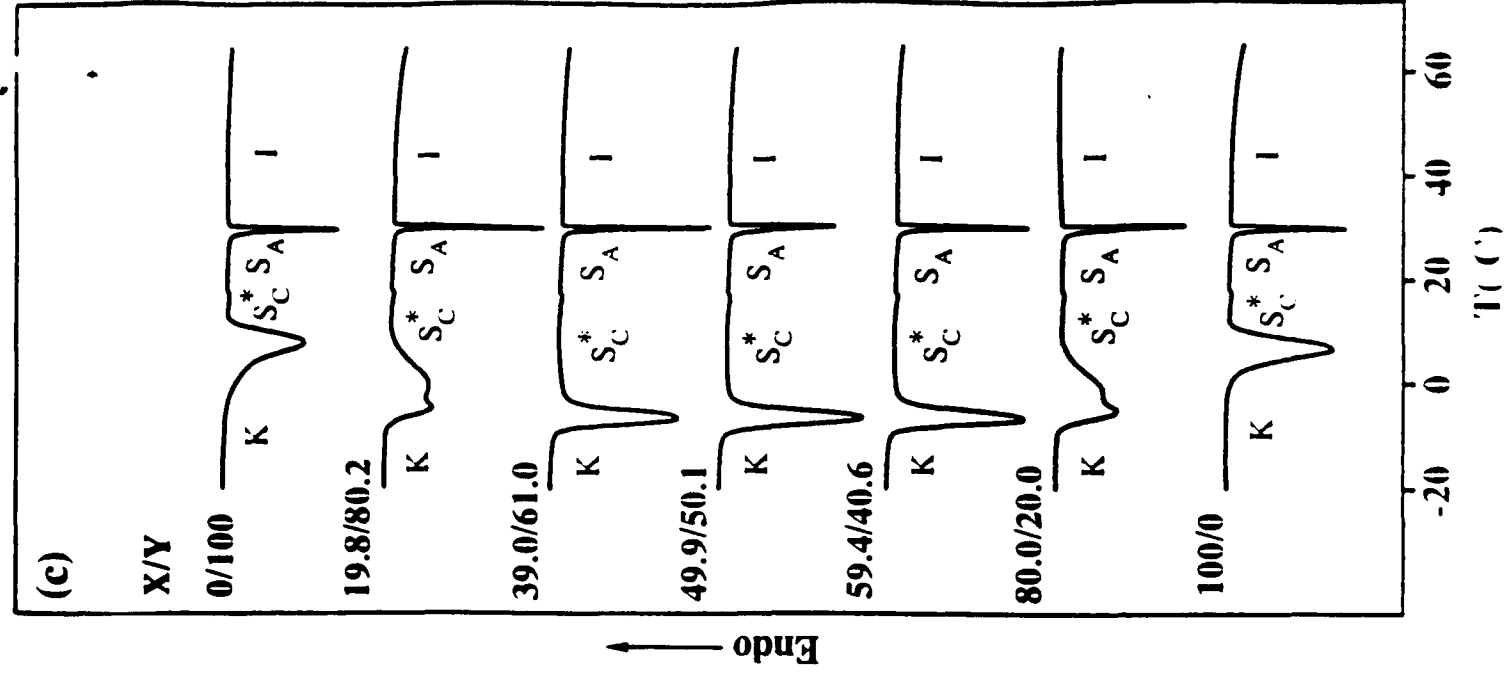
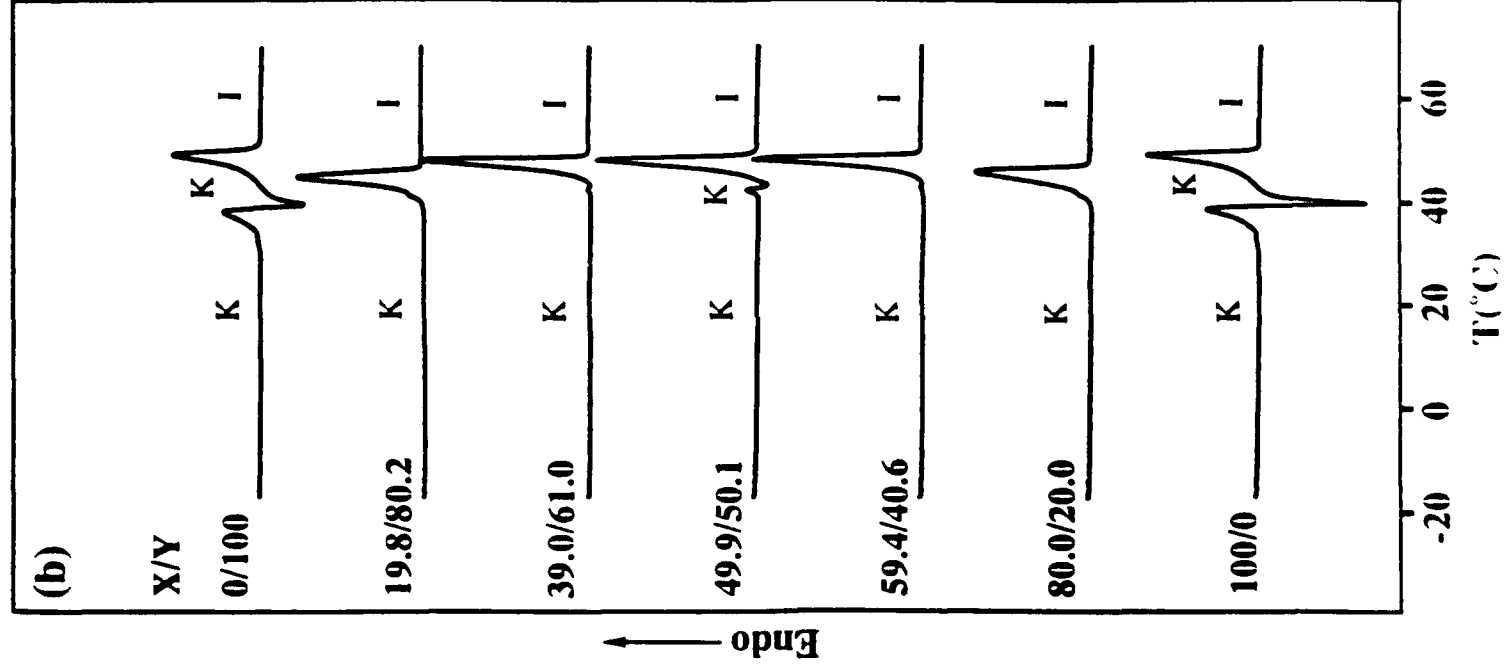
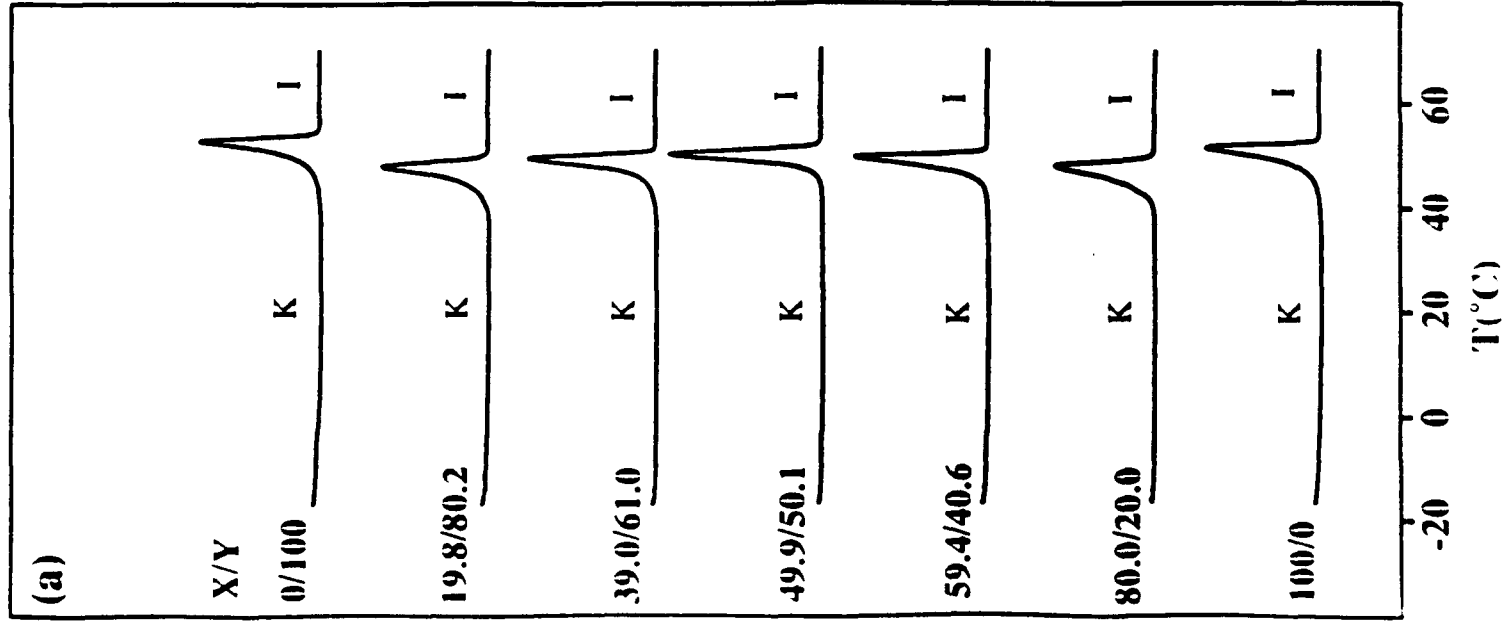
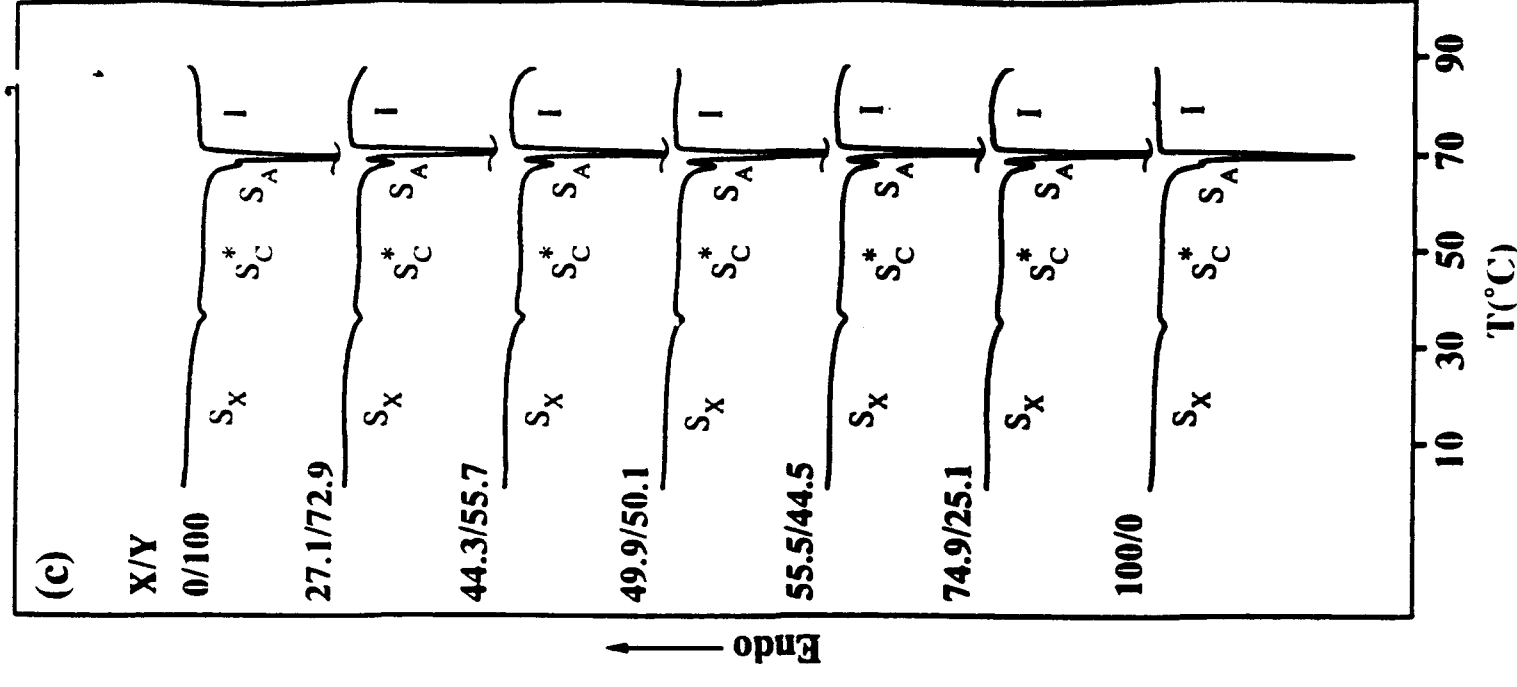
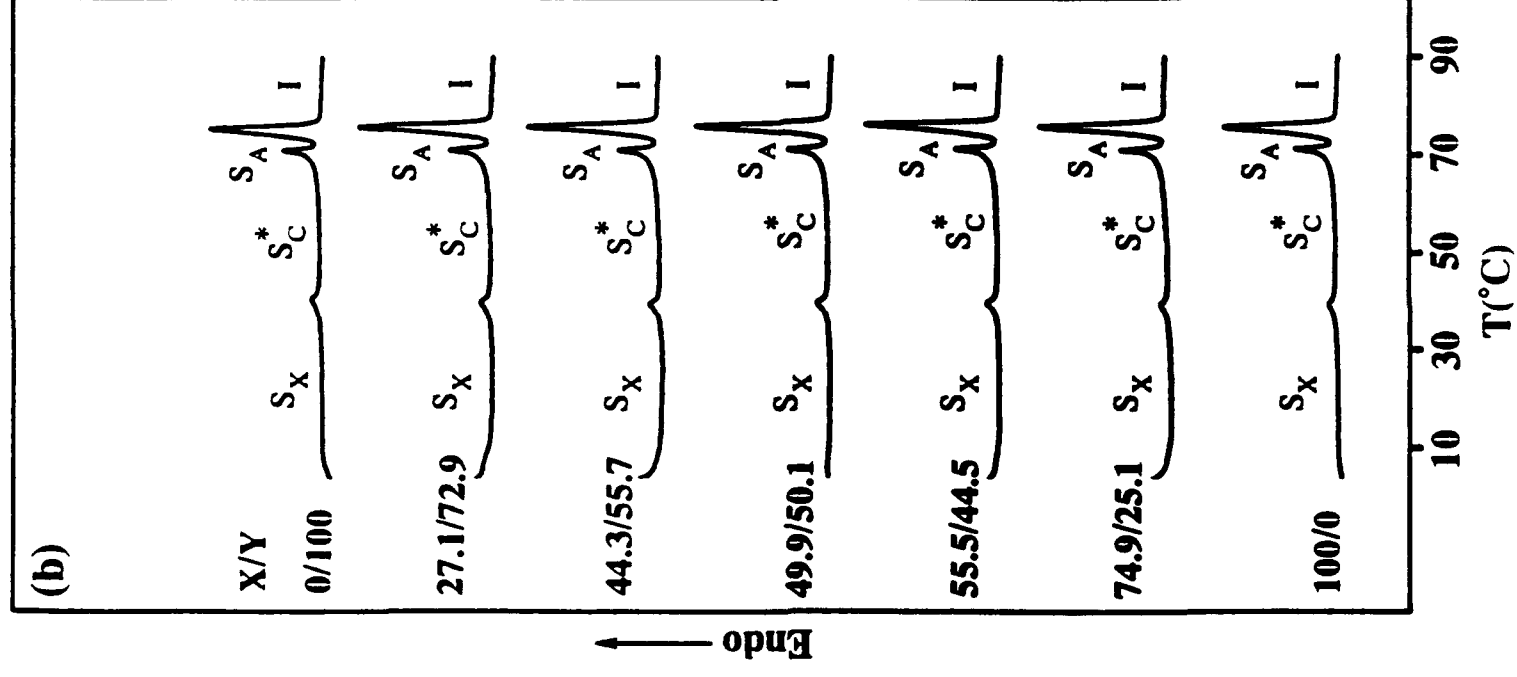
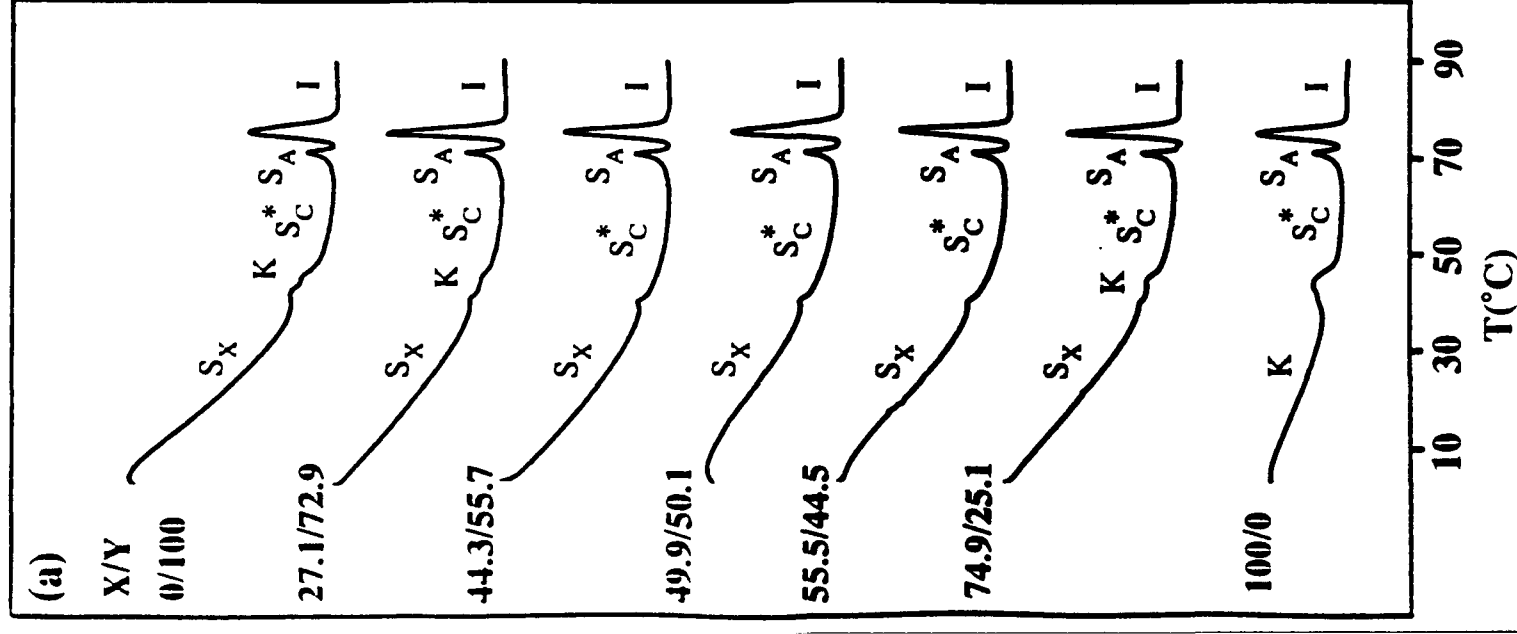


Figure 14





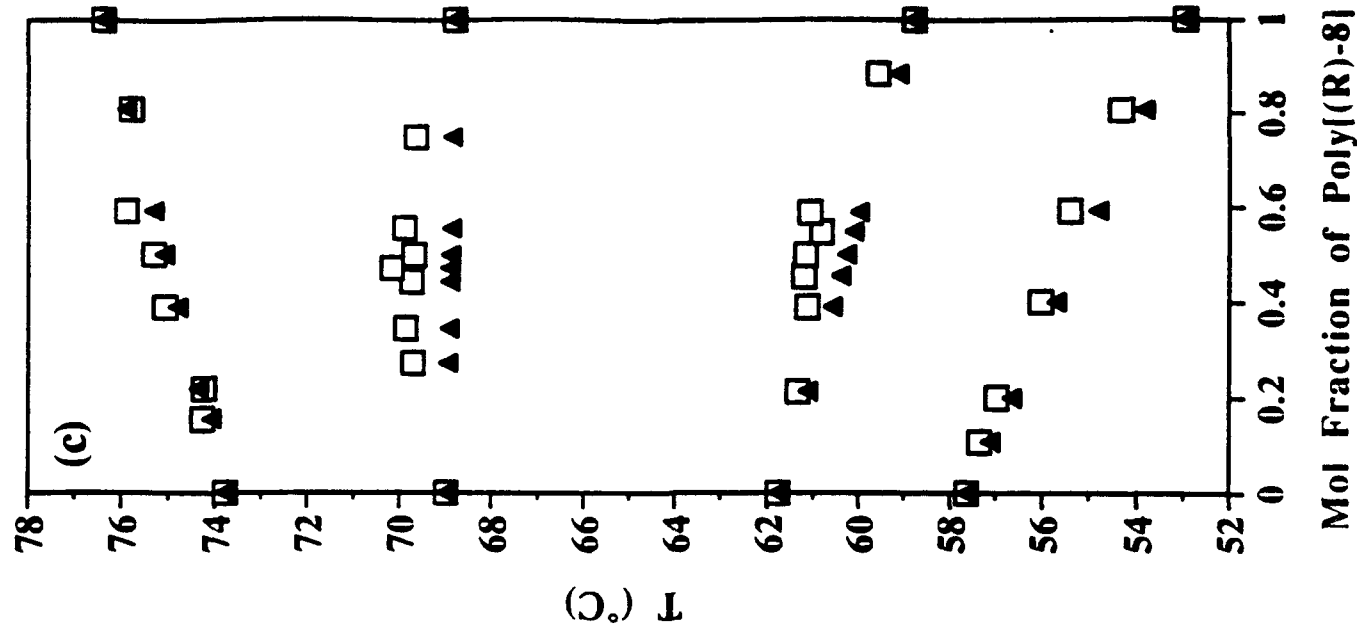
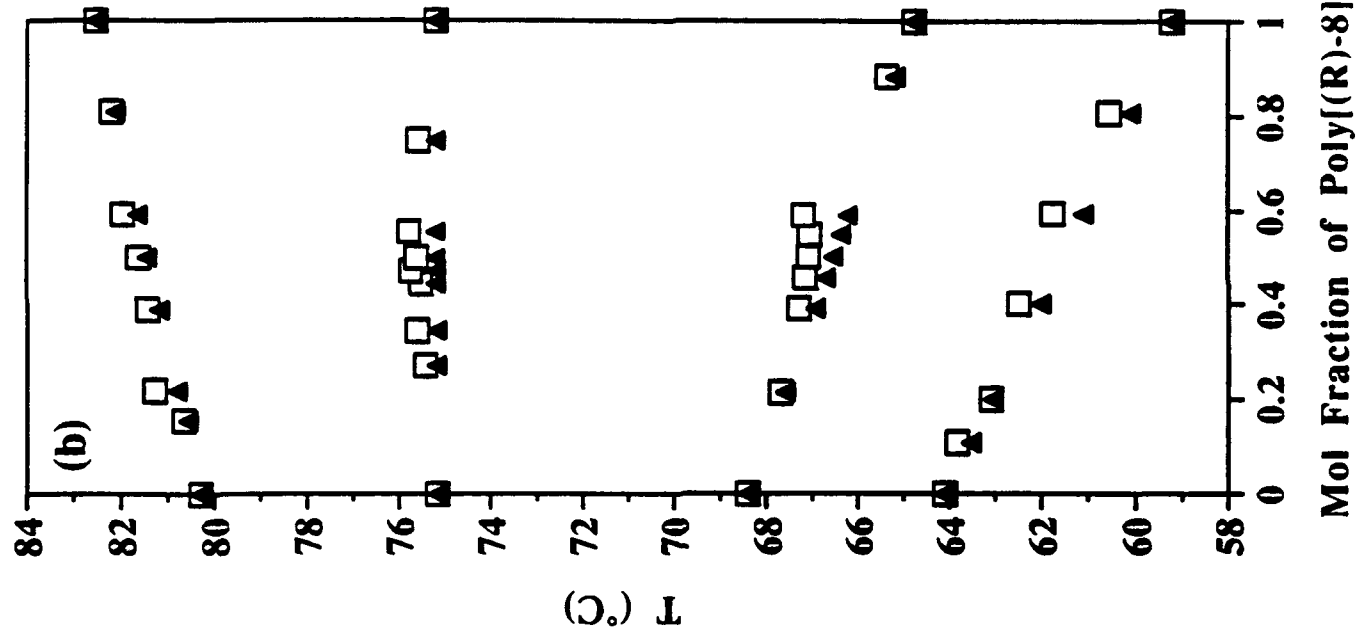
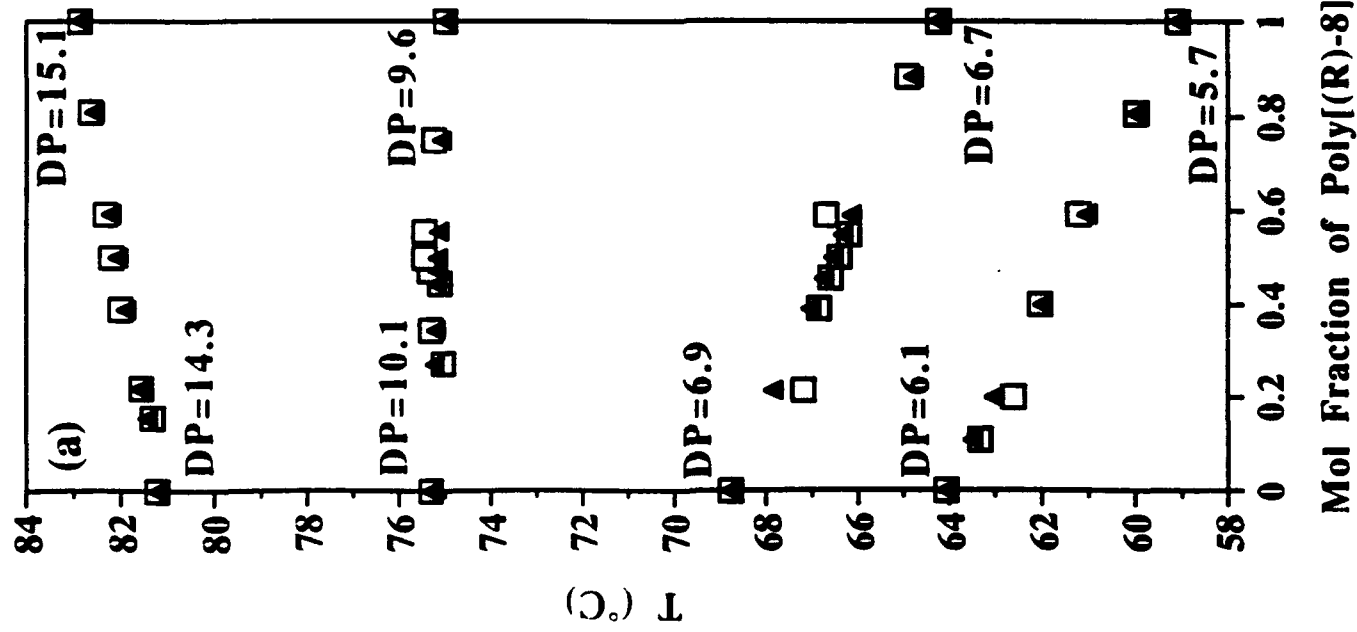
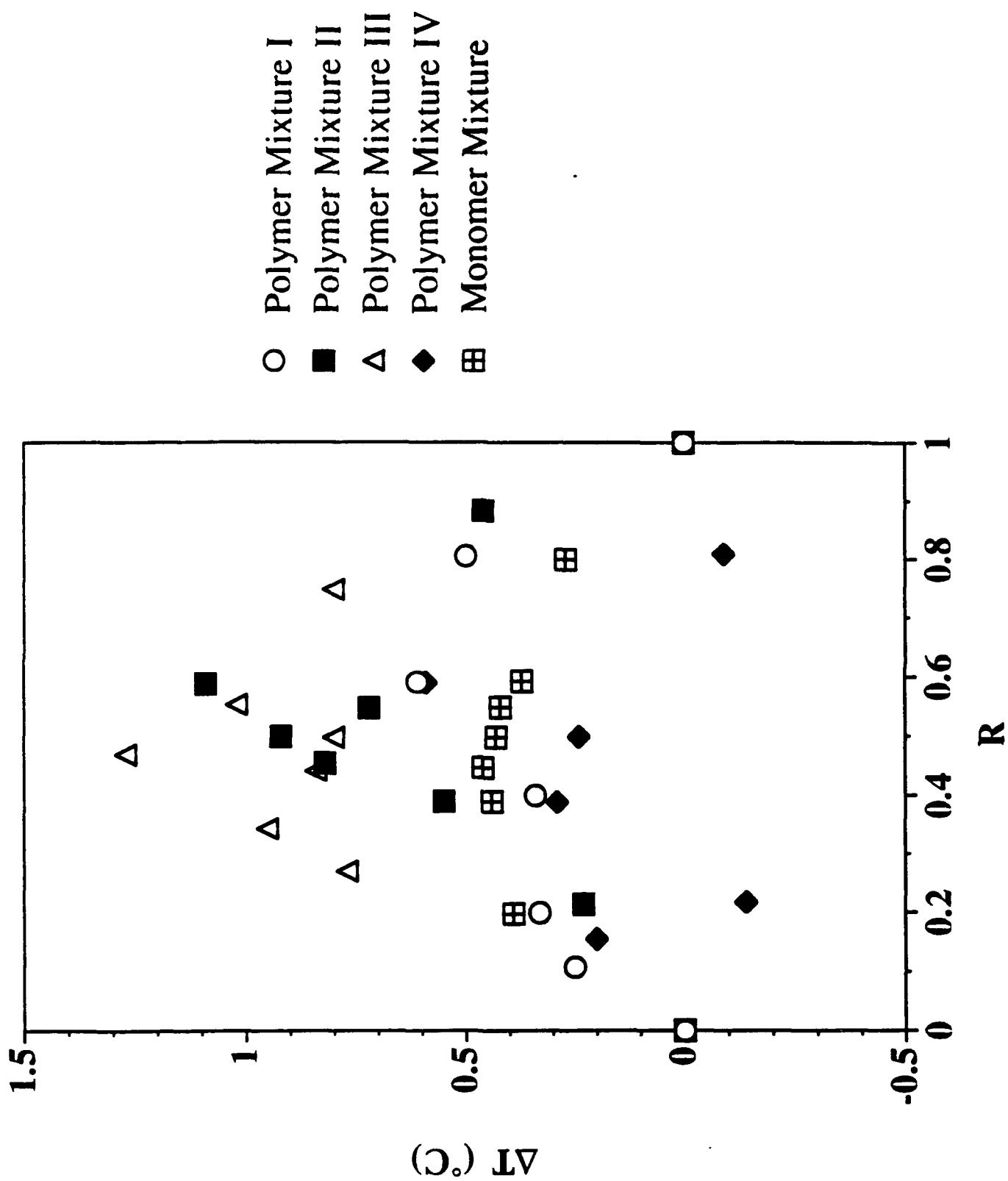


Figure 17

Figure 18



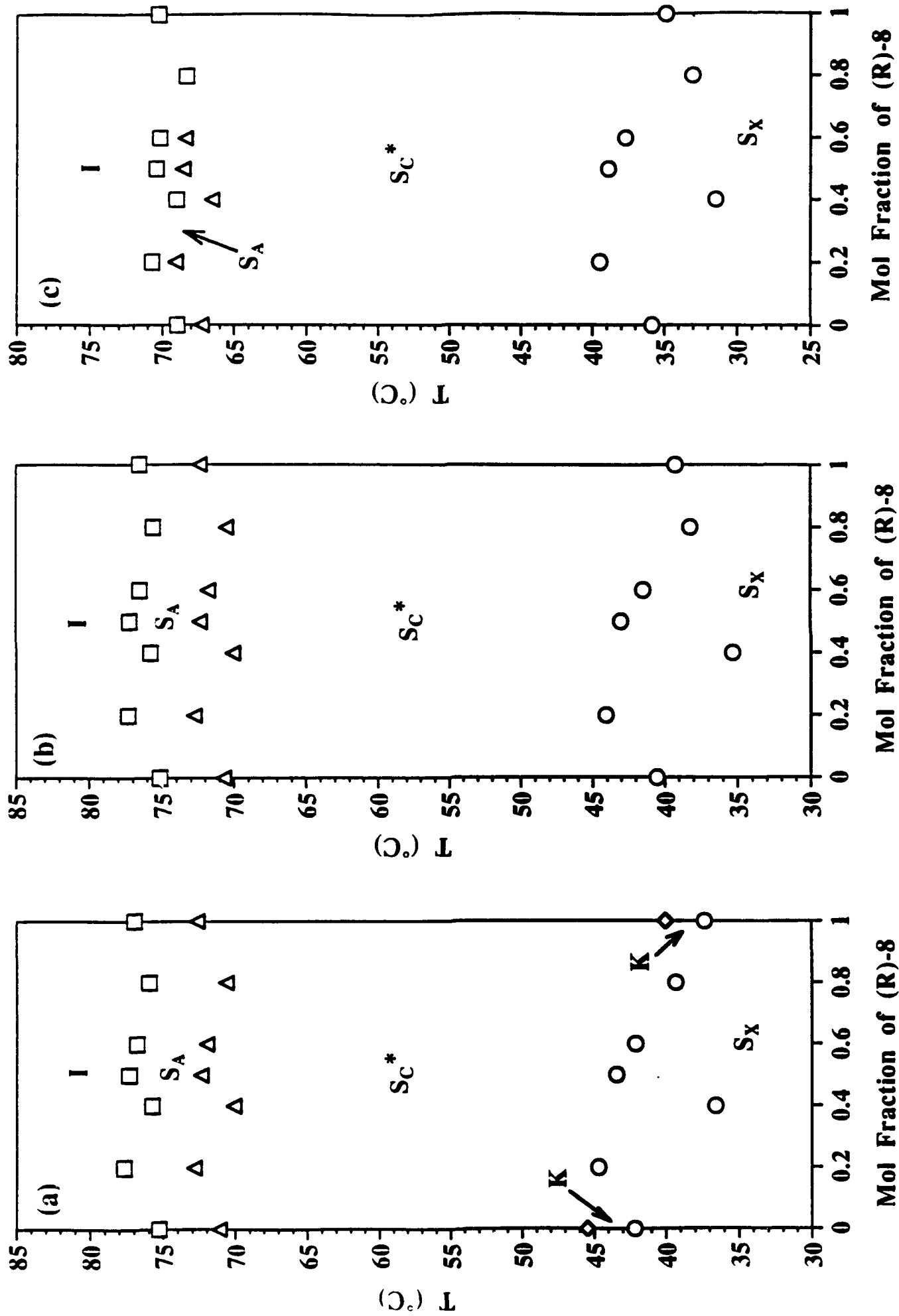


Figure 19